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(71) Applicant (for all designated States except US): **SYNGENTA PARTICIPATIONS AG** [CH/CH]; Schwarzwaldallee 215, 4058 Basel (CH).

(72) Inventor; and

(75) Inventor/Applicant (for US only): **TOBLER, Hans** [CH/CH]; Syngenta Crop Protection AG, Schwarzwaldallee 215, 4058 Basel (CH).

(74) Agent: **BASTIAN, Werner**; Syngenta Participations AG, Intellectual Property, P.O. Box, CH-4002 Basel (CH).

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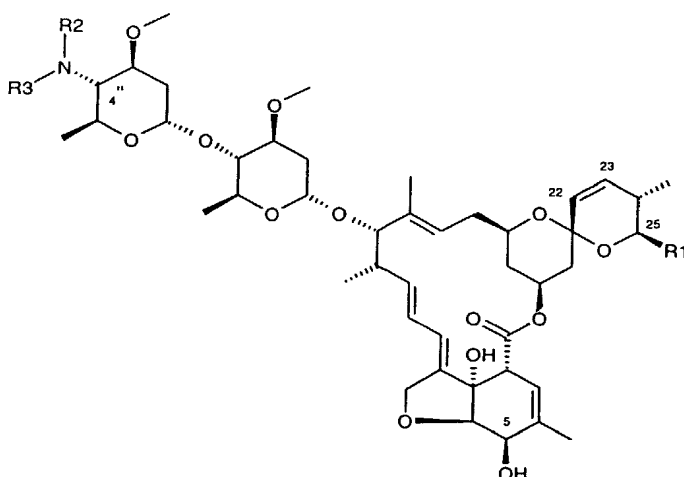
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(54) Title: 4"-DEOXY-4"-(S)-AMINO AVERMECTIN DERIVATIVES



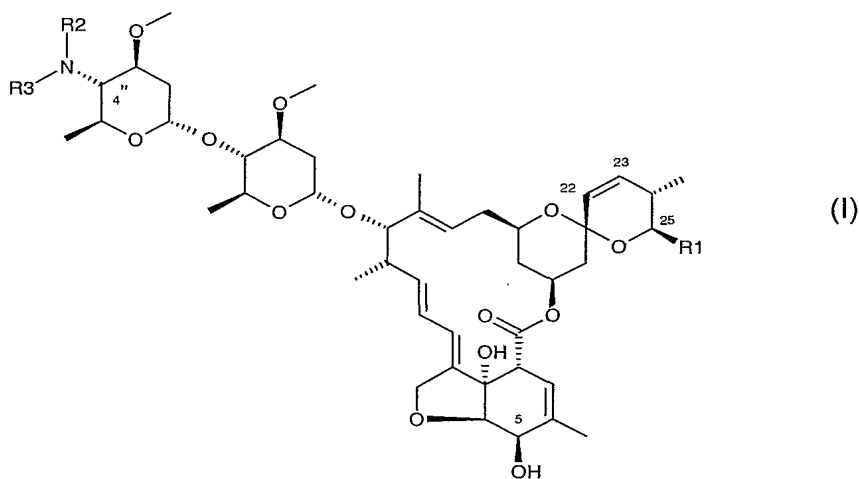
(I)

(57) Abstract: A description is given of compounds of the formula which in the 4"-position has the (S)-configuration and wherein R₁ is C₁-C₁₂alkyl, C₃-C₈cycloalkyl; or C₂-C₁₂alkenyl; R₂ is hydrogen, optionally substituted C₁-C₁₂alkyl or optionally substituted C₂-C₁₂alkenyl; R₃ is C₁-C₁₂alkyl, optionally substituted C₁-C₁₂alkyl, optionally substituted C₁-C₁₂alkoxy-C₁-C₁₂alkyl, optionally substituted C₃-C₁₂cycloalkyl, optionally substituted C₂-C₁₂alkenyl; optionally substituted C₄-C₁₂cycloalkenyl, optionally substituted C₂-C₁₂alkynyl; or R₂ and R₃ together are a three- to seven-membered alkylene or four-

to seven-membered alkenylene bridge in each of which a CH₂ group may have been replaced by O, S or NR₄; X is O or S; R₄ is C₁-C₈alkyl, C₃-C₈cycloalkyl, C₂-C₈alkenyl, C₂-C₈alkynyl, benzyl or C(=O)-R₅; R₅ is for example H, OH, SH, C₁-C₁₂alkyl, C₂-C₈alkenyl, C₂-C₈alkynylyl or C₁-C₁₂haloalkyl; and, where appropriate, E/Z isomers, mixtures of E/Z isomers and/or tautomers thereof, in each case in free form or in salt form; to a process for the preparation of, and to the use of, such compounds and isomers and tautomers thereof; to starting compounds for the preparation of compounds of formula (I); to pesticidal compositions in which the active ingredient has been selected from the compounds of formula (I) and tautomers thereof; and to a method of controlling pests using such compositions.

4"-Deoxy-4"-(S)-amino avermectin derivatives

The invention relates to (1) a compound of formula



which in the 4''-position has the (S)-configuration and wherein

R₁ is C₁-C₁₂alkyl, C₃-C₈cycloalkyl; or C₂-C₁₂alkenyl;

R₂ is hydrogen, unsubstituted or mono- to penta-substituted C₁-C₁₂alkyl or unsubstituted or mono- to penta-substituted C₂-C₁₂alkenyl;

R₃ is C₁-C₁₂alkyl, mono- to penta-substituted C₁-C₁₂alkyl, unsubstituted or mono- to penta-substituted C₁-C₁₂alkoxy-C₁-C₁₂alkyl, unsubstituted or mono- to penta-substituted C₃-C₁₂cycloalkyl, unsubstituted or mono- to penta-substituted C₂-C₁₂alkenyl; unsubstituted or mono- to penta-substituted C₄-C₁₂cycloalkenyl, unsubstituted or mono- to penta-substituted C₂-C₁₂alkynyl; or

R₂ and R₃ together are a three- to seven-membered alkylene or four- to seven-membered alkenylene bridge in each of which a CH₂ group may have been replaced by O, S or NR₄;

wherein the substituents of the mentioned alkyl, alkoxy-alkyl, alkenyl, alkynyl, alkylene, alkenylene, cycloalkyl and cycloalkenyl radicals are selected from the group consisting of OH, halogen, halo-C₁-C₂alkyl, CN, SCN, NO₂, Si(C₁-C₁₂alkyl)₃, C₂-C₆alkynyl, C₃-C₈cycloalkyl that is unsubstituted or substituted by from one to three methyl groups, norbornylenyl, C₃-C₈cycloalkenyl that is unsubstituted or substituted by from one to three methyl groups; C₃-C₈halocycloalkyl, C₁-C₁₂alkoxy, C₁-C₆alkoxy-C₁-C₆alkoxy, C₃-C₈cycloalkoxy, C₁-C₁₂halo-

alkoxy, C₁-C₁₂alkylthio, C₃-C₈cycloalkylthio, C₁-C₁₂haloalkylthio, C₁-C₁₂alkylsulfinyl, C₃-C₈cycloalkylsulfinyl, C₁-C₁₂haloalkylsulfinyl, C₃-C₈halocycloalkylsulfinyl, C₁-C₁₂alkylsulfonyl, C₃-C₈cycloalkylsulfonyl, C₁-C₁₂haloalkylsulfonyl, C₃-C₈halocycloalkylsulfonyl, C₂-C₈alkenyl, C₂-C₈alkynyl, NH₂, NH(C₁-C₆alkyl), NH(hydroxy-C₁-C₆alkyl), N(C₁-C₆alkyl)(hydroxy-C₁-C₆alkyl), N(phenyl)(hydroxy-C₁-C₆alkyl), N(C₁-C₆alkyl)₂, -C(=X)ZR₅, -O-C(=X)R₆, -O-C(=X)N(R₈)R₆, -NHC(=X)R₆, -NHC(=X)OR₆, -NHC(=X)SR₆, -NHC(=X)N(R₈)R₆, -S-C(=S)R₆, -NHS(O)₂-R₉, -P(=O)(OC₁-C₆alkyl)₂;

aryl, heterocyclyl, aryloxy, arylthio, heterocyclyloxy; and aryl, heterocyclyl, aryloxy, arylthio and heterocyclyloxy which, depending on the possibilities for substitution on the ring, are mono- to penta-substituted by substituents selected from the group consisting of OH, =O, halogen, CN, NO₂, C₁-C₁₂alkyl, hydroxyC₁-C₁₂alkyl, C₃-C₈cycloalkyl, C₁-C₁₂haloalkyl, C₁-C₁₂alkoxy, C₁-C₁₂haloalkoxy, C₁-C₁₂alkylthio, C₁-C₁₂haloalkylthio, C₁-C₆alkoxy-C₁-C₆alkyl, dimethylamino-C₁-C₆alkoxy, C₂-C₈alkenyl, C₂-C₈alkynyl, Si(C₁-C₁₂alkyl)₃, methylenedioxy, -C(=X)R₅, -O-C(=X)R₆, -CH₂-C(=O)R₅, -CH₂-O-C(=O)R₆, -NH-C(=X)R₆, -S-C(=S)R₆, NH₂, NH(C₁-C₁₂alkyl), -N(C₁-C₁₂alkyl)₂, C₁-C₆alkylsulfinyl, C₃-C₈cycloalkylsulfinyl, C₁-C₆haloalkylsulfinyl, C₃-C₈halocycloalkylsulfinyl, C₁-C₆alkylsulfonyl, C₃-C₈cycloalkylsulfonyl, C₁-C₆haloalkylsulfonyl and C₃-C₈halocycloalkylsulfonyl; unsubstituted phenyl, phenoxy, phenyl-C₁-C₆alkyl, phenyl-C₁-C₆alkoxy, phenyl-C₂-C₆alkenyl, phenyl-C₂-C₆alkynyl; phenyl, phenoxy, phenyl-C₁-C₆alkyl, phenyl-C₁-C₆alkoxy, phenyl-C₂-C₆alkenyl and phenyl-C₂-C₆alkynyl, each of which is substituted in the phenylring by from one to three substituents independently selected of one another from nitro, cyano, halogen, C₁-C₁₂alkoxy, haloC₁-C₁₂alkyl and haloC₁-C₁₂alkoxy;

X is O or S;

Z is a bond, O, NR₈ or S;

R₄ is C₁-C₈alkyl, C₃-C₈cycloalkyl, C₂-C₈alkenyl, C₂-C₈alkynyl, benzyl or -C(=O)-R₅;

R₅ is H, OH, SH, C₁-C₁₂alkyl, C₂-C₈alkenyl, C₂-C₈alkynylyl, C₁-C₁₂haloalkyl, C₂-C₈haloalkenyl, C₁-C₆alkoxy-C₁-C₆alkyl, -C₁-C₆S(O)₂-R₉, C₁-C₆alkyl which is mono- or disubstituted with substituents selected from COOH, -C(=O)O-CH₂-phenyl, unsubstituted or substituted phenyl, unsubstituted or substituted biphenyl, wherein the substituents are selected from cyano, halogen, nitro, trifluoromethyl, trifluoromethyl and benzyloxy; aryl, heterocyclyl, aryl-C₁-C₁₂alkyl, aryloxy-C₁-C₁₂alkyl, -C₁-C₆alkyl-C(=O)-R₇, -C₁-C₆alkyl-C(=O)-R₇; or aryl, heterocyclyl, aryl-C₁-C₁₂alkyl, aryloxy-C₁-C₁₂alkyl, each of which is substituted on the aromatic ring by from one to three substituents selected

independently of one another from halogen, nitro, cyano, C₁-C₆alkoxy, C₁-C₆haloalkyl and C₁-C₆haloalkoxy;

R₆ is H, C₁-C₂₄alkyl, C₁-C₆alkoxy-C₁-C₆alkyl, C₁-C₁₂hydroxyalkyl, C₁-C₁₂haloalkyl, C₂-C₈alkenyl, C₂-C₈alkynyl, C₃-C₈cycloalkyl, -C₁-C₆S(O)₂-R₉, aryl or phenyl-C₁-C₆alkyl; wherein the phenyl- and aryl-radicals may carry from one to three substituents selected independently of one another from nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl and C₁-C₆haloalkoxy;

R₇ is H, OH, C₁-C₁₂alkyl, C₁-C₁₂alkoxy, C₁-C₆alkoxy-C₁-C₆alkoxy, C₂-C₈alkenyloxy, phenyl, phenoxy, benzyloxy, NH₂, NH(C₁-C₁₂alkyl), N(C₁-C₁₂alkyl)₂, -NH-phenyl or -N(C₁-C₁₂alkyl)-phenyl;

R₈ is H, C₁-C₁₂alkyl, phenyl or benzyl; and

R₉ H, C₁-C₁₂alkyl, aryl, aryl-C₁-C₁₂alkyl, or aryl or aryl-C₁-C₁₂alkyl wherein the aryl radical may carry from one to three substituents selected independently of one another from nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl and C₁-C₆haloalkoxy;

and, where appropriate, E/Z isomers, mixtures of E/Z isomers and/or tautomers thereof, in each case in free form or in salt form;

to a process for the preparation of, and to the use of, such compounds and isomers and tautomers thereof; to starting compounds for the preparation of compounds of formula (I); to pesticidal compositions in which the active ingredient has been selected from the compounds of formula (I) and tautomers thereof; and to a method of controlling pests using such compositions.

Certain macrolide compounds are proposed for pest control in the literature. The biological properties of those known compounds are not entirely satisfactory, however, for which reason there is a need to provide further compounds having pesticidal properties, especially for the control of insects and representatives of the order Acarina. That problem is solved, in accordance with the invention, by the provision of the present compounds of formula (I), wherein the (S)-configuration applies in the 4"-position.

The compounds claimed in accordance with the invention are derivatives of avermectin. Avermectins are known to the person skilled in the art. They are a group of pesticidally active compounds, closely related to one another structurally, that are obtained by fermentation of a strain of the *Streptomyces avermitilis* microorganism. Derivatives of avermectins can be obtained *via* conventional chemical syntheses.

The avermectins obtainable from *Streptomyces avermitilis* are designated A1a, A1b, A2a, A2b, B1a, B1b, B2a and B2b. Compounds with the designation "A" have a methoxy radical in the 5-position; those compounds designated "B" have an OH group. The "a" series comprises compounds wherein the substituent R₁ (in position 25) is a sec-butyl radical; the "b" series have an isopropyl radical in the 25-position. The number 1 in the name of a compound indicates that atoms 22 and 23 are bonded by a double bond; the number 2 indicates that they are bonded by a single bond and carbon atom 23 carries an OH group. The above designations are retained in the description of the present invention, in order to indicate the specific structural type - in the case of the non-natural avermectin derivatives according to the invention - that corresponds to natural avermectin. In accordance with the invention there are claimed derivatives of compounds of the B1 series, more especially mixtures of derivatives of avermectin B1a and avermectin B1b wherein the (S)-configuration applies in the 4"-position.

Some of the compounds of formula (I) may be in the form of tautomers. Accordingly, any reference to the compounds of formula (I) hereinbefore and hereinafter is to be understood, where applicable, as including also corresponding tautomers, even if the latter are not specifically mentioned in every case.

The compounds of formula (I) and, where appropriate, tautomers thereof are capable of forming salts, for example acid addition salts. Those acid addition salts are formed, for example, with strong inorganic acids, such as mineral acids, e.g. sulfuric acid, a phosphoric acid or a hydrohalic acid, with strong organic carboxylic acids, such as unsubstituted or substituted, e.g. halo-substituted, C₁-C₄alkanecarboxylic acids, saturated or unsaturated dicarboxylic acids, hydroxycarboxylic acids, or with organic sulfonic acids, such as unsubstituted or substituted, e.g. halo-substituted, C₁-C₄alkane- or aryl-sulfonic acids. Furthermore, compounds of formula (I) having at least one acid group are capable of forming salts with bases. Suitable salts with bases are, for example, metal salts, such as alkali metal and alkaline earth metal salts, e.g. sodium, potassium and magnesium salts, and salts with ammonia or an organic amine, such as morpholine, piperidine, pyrrolidine, a mono-, di- or tri-lower alkylamine, e.g. ethyl-, diethyl-, triethyl- or dimethyl-propyl-amine, or a mono-, di- or tri-hydroxy-lower alkylamine, e.g. mono-, di- or tri-ethanolamine. In addition, corresponding internal salts may, where appropriate, also be formed. Preference is given, firstly, to the free form. Among the salts of compounds of formula (I) preference is given to agrochemically advantageous salts. The free compounds of formula (I) and salts thereof are to be understood hereinbefore and hereinafter as including, where appropriate, both the

corresponding salts and the free compounds of formula (I), respectively. The same is correspondingly true for tautomers of compounds of formula (I) and salts thereof.

Preferred anions of salts of compounds of formula (I) are the anion of a mineral acid, for example of sulfuric acid, a phosphoric acid or a hydrohalic acid;

the anion of an organic carboxylic acid, for example an unsubstituted or substituted, e.g. halo-substituted, C₁-C₄alkanecarboxylic acid, a saturated or unsaturated dicarboxylic acid, or a hydroxycarboxylic acid,

the anion of an organic sulfonic acid, for example an unsubstituted or substituted, e.g. halo-substituted, C₁-C₄alkane- or aryl-sulfonic acid; or

the anion of an active-H-C compound. Such active-H-C compounds especially include organic compounds that carry strongly electron-withdrawing substituents, for example nitriles, carbonyls or nitro groups. Preference is given to anions of compounds of formula Y₁-CH₂-Y₂ wherein Y₁ and Y₂ denote an electron-withdrawing group, with special preference being given to the anions of malodinitrile, cyanoacetic acid, esters of cyanoacetic acid, amides of cyanoacetic acid, acetoacetic acid, esters of acetoacetic acid, acetylacetone, cyanoacetone and barbituric acid; or

the anion of an acidic phenol, for example picric acid.

Very special preference is given to 1:1 salts of compounds of formula (I) with the following acids: benzoic acid, maleic acid, fumaric acid, 2-hydroxybenzoic acid, salicylic acid, thiosalicylic acid, malic acid, benzenesulfonic acid, barbituric acid, 2-ethylbutyric acid, thiomalic acid, 3,5-dihydroxy-benzoic acid, trimesic acid, D-(-)-quinic acid, 2-bromo-benzoic acid, 2-phenyl-benzoic acid, 3,3'-thiodipropionic acid, naphthalene-1-carboxylic acid, 5-sulfosalicylic acid, 2-methoxy-phenylacetic acid, benzene-1,2,4-tricarboxylic acid, 3-hydroxybenzoic acid, D-gluconic acid, 4,5-dichloro-phthalic acid, n-hexanoic acid (caproic acid), n-heptanoic acid (oenanthic acid), n-octanoic acid (caprylic acid), stearic acid, palmitic acid, 2,2'-dihydroxy-1,1'-dinaphthylmethane-3,3'-4,4'-methylene-bis(3-hydroxy-2-naphthoic acid), embonic acid, 4-methoxy-phenylacetic acid (homoanisic acid), 2-anisic acid (2-methoxybenzoic acid), adamantane-1-carboxylic acid, pyridine-3,4-dicarboxylic acid, 3,4-dihydroxybenzoic acid, 1-hydroxy-2-naphthoic acid (1-naphthol-2-carboxylic acid), 2,2'-oxydiacetic acid (diglycolic acid), O-ethyl-glycolic acid, (2-naphthylthio)-acetic acid (S-(2-naphthyl)-thioglycolic acid), 2-naphthoxy-acetic acid, perfluoro-octanoic acid, p-toluic acid, cyclohexanepropionic acid, 2,6-dihydroxypyridine-4-carboxylic acid (citrazinic acid), 3-methoxypropionic acid,

3,4,5-trihydroxy-benzoic acid (gallic acid), pyromucic acid (furan-2-carboxylic acid), 2-methylbenzoic acid (o-toluic acid), 3,6,9-trioxa-undecanedioic acid, 3-(4-methoxyphenyl)-propionic acid (p-methoxy-hydrocinnamic acid), 3-(3,4-dihydroxyphenyl)-propionic acid, O-acetyl-salicylic acid (aspirin), 3-fluoro-benzoic acid, cyclohexanecarboxylic acid, 5-chloro-2-hydroxybenzoic acid (5-chloro-salicylic acid), 2,5-dimethyl-benzoic acid (p-xylylic acid), 3,4,5-trimethoxy-benzoic acid (trimethylgallic acid), 2,4,6-trimethyl-benzoic acid, 3-phenoxy-benzoic acid, 4-phenyl-butyric acid, 3-trifluoromethyl-benzoic acid, terephthalic acid monomethyl ester, o-hydroxy-phenyl-acetic acid, isophthalic acid, 2,4,6-trihydroxy-benzoic acid, trifluoromethanesulfonic acid, 2-methyl-propionic acid (isobutyric acid), 2,4-dimethoxy-benzoic acid, 2-thienylacetic acid (thiophene-2-acetic acid), 3,4-dimethoxy-benzoic acid (veratric acid), 2,2-bis(hydroxymethyl)-propionic acid, 2-fluoro-phenylacetic acid, 2-methyl-butyric acid, hydroxy-acetic acid, 4-chloro-phenylacetic acid, 2-mercaptobenzoic acid (thiosalicylic acid), (+/-)-2-hydroxyphenyl-acetic acid (DL-mandelic acid), 2,4-dihydroxypyrimidine-6-carboxylic acid, toluene-4-sulfonic acid, (p-toluene-sulfonic acid), 2-chloro-phenylacetic acid, 2,4-dichloro-benzoic acid, 2,6-dichloro-benzoic acid, 2-mercapto-propionic acid (thiolactic acid), 2-chloro-benzoic acid, methanesulfonic acid, ethanesulfonic acid (ethyl-sulfuric acid), 4-phenoxy-butyric acid, 4-tert-butyl-benzoic acid, 3,4-methylenedioxy-benzoic acid, bis(2-carboxyethyl)-disulfide, pivalic acid (trimethylacetic acid), nicotinic acid N-oxide, acrylic acid, 3-benzoylpropionic acid (4-oxo-4-phenyl-butyric acid), (1R)-(-)-camphor-10-sulfonic acid hydrate, 2-chloro-4-fluoro-benzoic acid, 3,5-dimethoxy-benzoic acid, 2-sulfobenzoic acid, sulfoacetic acid, 2-chloro-6-fluoro-benzoic acid, 2,4-dihydroxy-benzoic acid, methoxyacetic acid, 2,4,6-trimethyl-benzene-sulfonic acid, tartaric acid, xanthene-9-carboxylic acid, 4-pentenoic acid (allylacetic acid), 5-sulfosalicylic acid, vinylacetic acid, 2-butyneedioic acid (acetylenedicarboxylic acid), 2-oxo-propionic acid (pyruvic acid), cyclohexylacetic acid, 2-hydroxyisobutyric acid, nicotinic acid, 6-chloro-nicotinic acid, isonicotinic acid, picolinic acid, pyrazinecarboxylic acid, oxalic acid, propionic acid, pentafluoropropionic acid, butyric acid, heptafluorobutyric acid, valeric acid, citric acid, glyceric acid, acetic acid, chloroacetic acid, dichloroacetic acid, trifluoroacetic acid, fluoroacetic acid, lactic acid, malonic acid, succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelaic acid, sebacic acid, phthalic acid, terephthalic acid, phosphoric acid, sulfuric acid, hydrochloric acid, hydrobromic acid, hydriodic acid, nitric acid, perchloric acid, acetoacetic acid, cyanoacetic acid, tetrahydrofuran-2-carboxylic acid, propiolic acid, methacrylic acid, crotonic acid and picric acid.

The general terms used hereinbefore and hereinafter have the following meanings, unless defined otherwise.

Unless defined otherwise, carbon-containing groups and compounds each contain from 1 up to and including 6, preferably from 1 up to and including 4, especially 1 or 2, carbon atoms.

Halogen, both as a group *per se* and as a structural element of other groups and compounds, for example haloalkyl, haloalkoxy and haloalkylthio, is fluorine, chlorine, bromine or iodine, especially fluorine, chlorine or bromine, more especially fluorine or chlorine.

Alkyl, both as a group *per se* and as a structural element of other groups and compounds, for example haloalkyl, alkoxy and alkylthio, is, in each case taking due account of the particular number of carbon atoms contained in the group or compound in question, either straight-chained, i.e. methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl or octyl, or branched, e.g. isopropyl, isobutyl, sec-butyl, tert-butyl, isopentyl, neopentyl or isohexyl.

Cycloalkyl, both as a group *per se* and as a structural element of other groups and compounds, for example halocycloalkyl, cycloalkoxy and cycloalkylthio, is, in each case taking due account of the particular number of carbon atoms contained in the group or compound in question, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or cyclooctyl.

Alkenyl, both as a group *per se* and as a structural element of other groups and compounds, is, in each case taking due account of the particular number of carbon atoms and conjugated or isolated double bonds contained in the group in question, either straight-chained, for example vinyl, allyl, 2-butenyl, 3-pentenyl, 1-hexenyl, 1-heptenyl, 1,3-hexadienyl or 1,3-octadienyl, or branched, for example isopropenyl, isobutenyl, isoprenyl, tert-pentenyl, isohexenyl, isoheptenyl or isooctenyl. Preference is given to alkenyl groups containing from 3 to 12, especially from 3 to 6, more especially 3 or 4, carbon atoms.

Alkynyl, both as a group *per se* and as a structural element of other groups and compounds, is, in each case taking due account of the particular number of carbon atoms and conjugated or isolated double bonds contained in the group or compound in question, either straight-chained, for example ethynyl, propargyl, 2-butyne, 3-pentyne, 1-hexynyl, 1-heptynyl, 3-hexen-1-ynyl or 1,5-heptadien-3-ynyl, or branched, for example 3-methylbut-1-ynyl, 4-ethylpent-1-ynyl, 4-methylhex-2-ynyl or 2-methylhept-3-ynyl. Preference is given to alkynyl groups containing from 3 to 12, especially from 3 to 6, more especially 3 or 4, carbon atoms.

Alkylene and alkenylene are straight-chained or branched bridge elements, especially $-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, $-\text{CH}_2(\text{CH}_3)\text{CH}_2-\text{CH}_2-$, $-\text{CH}_2\text{C}(\text{CH}_3)_2-\text{CH}_2-$, $-\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2-$ or $-\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2-\text{CH}_2-$.

Halo-substituted, carbon-containing groups and compounds, for example halo-substituted alkyl, alkenyl, alkynyl, cycloalkyl, alkoxy and alkylthio, may be partially halogenated or perhalogenated, the halogen substituents in the case of multiple halogenation being identical or different. Examples of haloalkyl, both as a group *per se* and as a structural element of other groups and compounds, for example haloalkoxy and haloalkylthio, are methyl mono- to tri-substituted by fluorine, chlorine and/or by bromine, for example CHF_2 or CF_3 ; ethyl mono- to penta-substituted by fluorine, chlorine and/or by bromine, for example CH_2CF_3 , CF_2CF_3 , CF_2CCl_3 , CF_2CHCl_2 , CF_2CHF_2 , CF_2CFCl_2 , CF_2CHBr_2 , CF_2CHClF , CF_2CHBrF or CClFCHClF ; propyl or isopropyl each mono- to hepta-substituted by fluorine, chlorine and/or by bromine, for example $\text{CH}_2\text{CHBrCH}_2\text{Br}$, $\text{CF}_2\text{CHFCH}_2\text{F}$, $\text{CH}_2\text{CF}_2\text{CF}_3$ or $\text{CH}(\text{CF}_3)_2$; butyl, or one of the isomers thereof, mono- to nona-substituted by fluorine, chlorine and/or by bromine, for example $\text{CF}(\text{CF}_3)\text{CH}_2\text{CF}_3$ or $\text{CH}_2(\text{CF}_2)_2\text{CF}_3$; pentyl, or one of the isomers thereof, mono- to undeca-substituted by fluorine, chlorine and/or by bromine, for example $\text{CF}(\text{CF}_3)(\text{CHF})_2\text{CF}_3$ or $\text{CH}_2(\text{CF}_2)_3\text{CF}_3$; and hexyl, or one of the isomers thereof, mono- to trideca-substituted by fluorine, chlorine and/or by bromine, for example $(\text{CH}_2)_4\text{CHBrCH}_2\text{Br}$, $\text{CF}_2(\text{CHF})_4\text{CF}_3$, $\text{CH}_2(\text{CF}_2)_4\text{CF}_3$ or $\text{C}(\text{CF}_3)_2(\text{CHF})_2\text{CF}_3$.

Aryl is especially phenyl, naphthyl, anthryl or perylenyl, preferably phenyl.

Heterocyclyl is especially pyridyl, pyrimidyl, s-triazinyl, 1,2,4-triazinyl, thienyl, furyl, tetrahydrofuranlyl, pyranlyl, tetrahydropyranlyl, pyrrolyl, pyrazolyl, imidazolyl, thiazolyl, triazolyl, oxazolyl, thiadiazolyl, oxadiazolyl, benzothienyl, quinolyl, quinoxalinyl, benzofuranlyl, benzimidazolyl, benzopyrrolyl, benzothiazolyl, indolyl, tetrahydro-thiopyranlyl, benzopyranlyl, N-oxido-pyridinio, fluorenyl, thiomorpholin-yl, morpholinyl, piperidinyl, pyrrolidinyl, piperazinyl, dihydropyranlyl, dihydroindolyl, piperazinyl, coumarinyl or indazolyl which are preferably attached by way of a carbon atom; preference is given to thienyl, thiazolyl, benzofuranlyl, benzothiazolyl, furyl, oxiranyl, tetrahydropyranlyl or indolyl; especially pyridyl or thiazolyl.

In the context of the present invention, preference is given to

(2) compounds according to group (1) of formula (I) wherein R_1 is isopropyl or sec-butyl, preferably wherein there is present a mixture of the isopropyl and sec-butyl derivatives;

- (3) compounds according to group (2) of formula (I) wherein R_2 is H;
- (4) compounds according to group (2) of formula (I) wherein R_2 is C_1 - C_8 alkyl, especially methyl;
- (5) compounds according to group (2) of formula (I) wherein R_2 is ethyl;
- (6) compounds according to group (2) of formula (I) wherein R_2 is n-propyl;
- (7) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is unsubstituted or substituted, especially unsubstituted, C_1 - C_{12} alkyl;
- (8) compounds according to one of groups (1) to (7) of formula (I) wherein R_3 is methyl;
- (9) compounds according to one of groups (1) to (7) of formula (I) wherein R_3 is ethyl;
- (10) compounds according to one of groups (1) to (7) of formula (I) wherein R_3 is n-propyl;
- (11) compounds according to one of groups (2) to (7) of formula (I) wherein R_3 is isopropyl;
- (12) compounds according to one of groups (1) to (7) of formula (I) wherein R_3 is n-butyl, sec-butyl, isobutyl or tert-butyl;
- (13) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is unsubstituted or substituted, especially unsubstituted, C_6 - C_{12} alkyl;
- (14) compounds according to one of groups (1) and (2) of formula (I) wherein R_2 and R_3 together are $-CH_2-CH_2-CH_2-$ or $-CH_2-CH_2-CH_2-CH_2-$;
- (15)) compounds according to one of groups (1) and (2) of formula (I) wherein R_2 and R_3 together are $-CH_2-CH_2-O-CH_2-CH_2-$ or $-CH_2-CH_2-N(CH_3)-CH_2-CH_2-$;
- (16) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is substituted C_1 - C_4 alkyl and the substituents are selected from the group consisting of OH, halogen, C_3 - C_8 cycloalkyl, C_3 - C_8 cycloalkenyl that is unsubstituted or substituted by from one to three methyl groups, C_1 - C_{12} alkoxy, C_2 - C_8 alkynyl, $-C(=O)R_5$, $-NHC(=O)R_6$, $-P(=O)(OC_1-C_6alkyl)_2$, unsubstituted or, depending on the possibilities for substitution on the ring, mono- to penta-substituted phenyl, naphthyl, anthryl, phenanthrenyl, fluorenyl, perylenyl and heterocyclyl;

especially wherein the substituents of R_3 are selected from the group consisting of halogen, C_3 - C_8 cycloalkyl, C_2 - C_8 alkynyl, $-C(=O)R_5$, $-NHC(=O)R_6$, $-P(=O)(OC_1-C_6alkyl)_2$; phenyl, naphthyl, anthryl, pyridyl, thiazolyl, imidazolyl, furyl, quinolyl, pyrazolyl, which are unsubstituted or, depending on the possibilities for substitution on the ring, are mono- to tri-substituted;

(17) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is benzyl carrying, on the aromatic moiety, from one to three substituents selected from the group consisting of OH, halogen, CN, NO_2 , C_1 - C_2 alkyl, dimethylamino- C_1 - C_4 alkoxy, C_3 - C_6 cycloalkyl, C_1 - C_2 haloalkyl, C_1 - C_2 alkoxy, C_1 - C_2 haloalkoxy, phenoxy, phenyl- C_1 - C_6 alkyl, phenyl- C_1 - C_4 alkenyl, unsubstituted or chloro- or methoxy-substituted phenoxy, unsubstituted or chloro-, methoxy- or trifluoromethyl-substituted benzyloxy, methylenedioxy, $-C(=O)ZR_5$, $-O-C(=O)R_6$ and $NHC(=O)R_6$;

R_5 is H, OH, NH_2 , $NH(C_1-C_2alkyl)$, $N(C_1-C_2alkyl)_2$, $-O-C_1-C_2alkyl-C(=O)-R_7$, $NHC_1-C_2alkyl-C(=O)-R_7$, C_1-C_6alkyl , $C_1-C_2alkoxy$, $C_1-C_2alkoxy-C_1-C_2alkoxy$, $C_2-C_4alkenyloxy$, $C_2-C_4alkynyloxy$; phenyl, phenoxy, benzyloxy, NH-phenyl, $NH-C_1-C_6alkyl-C(=O)-R_7$; or phenyl, phenoxy, benzyloxy or NH-phenyl which are substituted by halogen, nitro, methoxy, trifluoromethyl or trifluoromethoxy;

R_6 is H, C_1 - C_3alkyl , phenyl or benzyl;

R_7 is H, OH, NH_2 , $NH(C_1-C_{12}alkyl)$, $N(C_1-C_{12}alkyl)_2$, $C_1-C_{12}alkyl$, $C_1-C_{12}alkoxy$, $C_1-C_6alkoxy-C_1-C_6alkoxy$, $C_2-C_8alkenyloxy$, phenyl, phenoxy, benzyloxy, or NH-phenyl; and

Z is a bond, O or NH;

(18) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is C_1 - $C_4alkyl-C(=O)R_5$, especially $-CH_2-C(=O)R_5$; and

R_5 is H, OH, NH_2 , $NH(C_1-C_2alkyl)$, $N(C_1-C_2alkyl)_2$, C_1-C_4alkyl , $C_1-C_{12}alkoxy$, $C_2-C_4alkenyloxy$, phenyl, phenoxy, benzyloxy, NH-phenyl, $NH-C_1-C_2alkyl-C(=O)-O-C_1-C_2alkyl-phenyl$, $-P(=O)(OC_1-C_6alkyl)_2$; or phenyl, phenoxy, benzyloxy or NH-phenyl which are substituted by chlorine, fluorine, methoxy, trifluoromethyl or by trifluoromethoxy;

more especially wherein R_5 is C_1 - $C_{12}alkoxy$;

(19) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is C_2 - $C_6alkyl-NHC(=O)R_6$ and R_6 is H, C_1 - C_4alkyl , C_2 - $C_4alkenyl$, C_2 - $C_4alkynyl$, phenyl or benzyl;

(20) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is $-\text{CH}_2$ -heterocyclyl and heterocyclyl is pyridyl, furyl, tetrahydrofuranyl, pyranyl, tetrahydropyranyl, pyrazolyl, imidazolyl, thiazolyl, benzothienyl, quinolyl, quinoxaliny, benzofuranyl, benzimidazolyl, benzopyrrolyl, benzothiazolyl, indolyl, coumarinyl or indazolyl, the mentioned radicals being unsubstituted or substituted by one or two substituents selected independently of one another from halogen, trifluoromethyl, trifluoromethoxy and nitro; preferably pyridyl, furyl, pyrazolyl, imidazolyl, thiazolyl, benzimidazolyl, benzopyrrolyl, benzothiazolyl or indolyl which are unsubstituted or substituted by one or two substituents selected independently of one another from halogen, trifluoromethyl, trifluoromethoxy and nitro; especially pyridyl or thiazolyl which are unsubstituted or substituted by one or two substituents selected independently of one another from halogen, trifluoromethyl, trifluoromethoxy and nitro, more especially monosubstituted by chlorine;

(21) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is C_2 - C_{10} alkenyl, especially C_2 - C_4 alkenyl, which is unsubstituted or mono- or di-substituted, especially mono-substituted, by C_2 - C_4 alkynyl, $-\text{C}(=\text{O})-\text{C}_1$ - C_4 alkoxy, $-\text{C}(=\text{O})-\text{O}-\text{C}_1$ - C_4 alkyl-benzoyl, phenyl or by halogen; especially wherein R_3 is $-\text{CH}_2-\text{CH}=\text{CH}_2$;

(22) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is branched, unsubstituted C_4 - C_{10} alkyl;

(23) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is branched, substituted C_3 - C_{10} alkyl;

(24) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is unsubstituted benzyl;

(25) compounds according to one of groups (1) to (6) of formula (I) wherein R_3 is unsubstituted or substituted, especially unsubstituted, C_2 - C_{12} alkyl;

(26) in the context of the invention special preference is given to the compounds of formula (I) listed in the Tables and, where appropriate, E/Z isomers and mixtures of E/Z isomers thereof, very especially the compounds

4"-deoxy-4"-(S)-N-methylamino-avermectin B1;

4"-deoxy-4"-(S)-N-dimethylamino-avermectin B1;

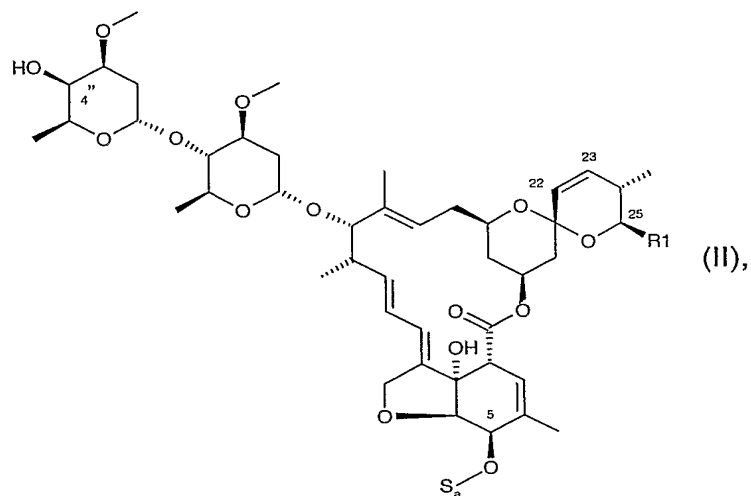
4"-deoxy-4"-(S)-N-ethylamino-avermectin B1;

4"-deoxy-4"-(S)-N-prop-1-ylamino-avermectin B1;

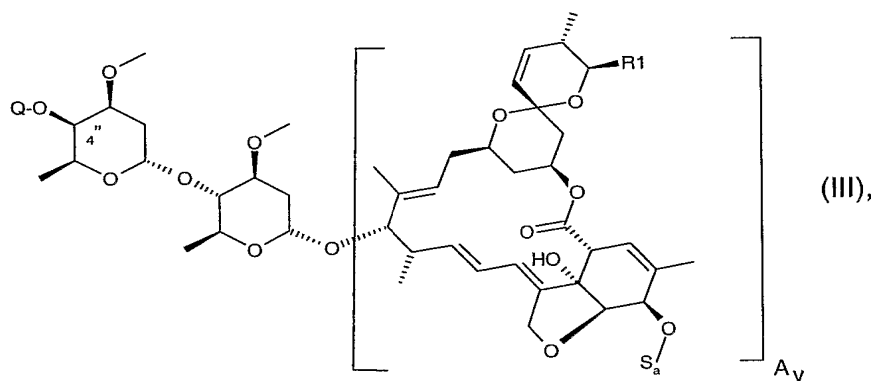
4"-deoxy-4"-(S)-(N-ethyl-N-methyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-prop-1-yl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-isopropyl-N-methylamino)-avermectin B1;
4"-deoxy-4"-(S)-(N-isopropyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-1-propen-3-yl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-ethoxycarbonylmethyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-benzyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-4-difluoromethoxyphenylmethyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-2,5-dichlorophenylmethyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-2,5-difluorophenylmethyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-2,3,4-trifluorophenylmethyl-amino)-avermectin B1;
4"-deoxy-4"-(S)-(pyrrolidin-1-yl)-avermectin B1;
4"-deoxy-4"-(S)-(azetidin-1-yl)-avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-[3-{2-oxo-2-phenyl-ethoxycarbonyl}-allyl]-amino)-
avermectin B1;
4"-deoxy-4"-(S)-(N-methyl-N-1-propargyl-amino)-avermectin B1; and
4"-deoxy-4"-(S)-(piperidin-1-yl)-avermectin B1.

The invention relates also to a process for the preparation of compounds of formula (I) as defined under (1) hereinbefore and, where appropriate, tautomers thereof, in which process

(A) a compound of formula

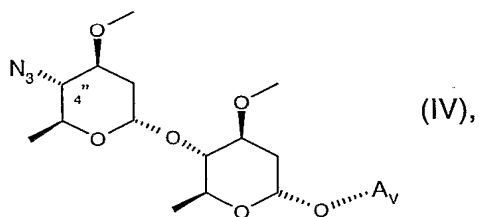


wherein R_1 is as defined hereinbefore under (1) for formula (I) and S_a is a protecting group, and which is known or can be prepared by methods known *per se*, is reacted with a sulfonic acid derivative to form a compound of formula



wherein Q is a sulfonic acid radical and S_a and R_1 are as defined for formula (II);

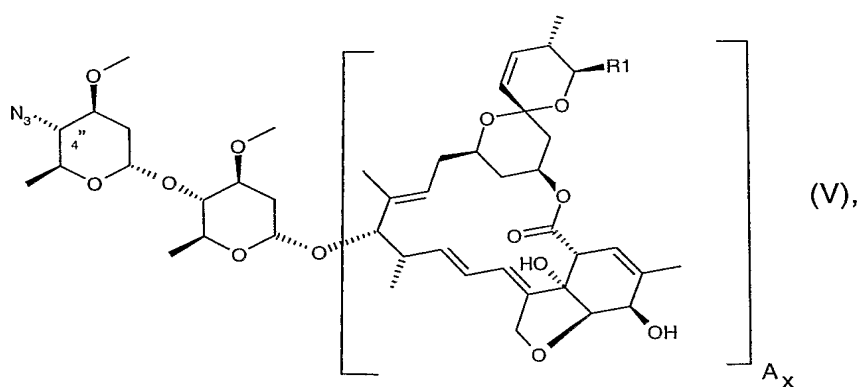
(B) the resulting compound of formula (III) is reacted with an azide salt to form a compound of formula



wherein the radical A_V corresponds to the macrocycle of the basic avermectin structure as indicated within the brackets of formula (III), with inversion at the 4"-position; or, where appropriate,

(C) for the preparation of a compound of formula (IV), a compound of formula (II) is reacted with an azide in the presence of triphenylphosphine or a trialkylphosphine and an azodicarboxylic acid derivative;

(D) the protecting group S_a of the compound of formula (IV) is split off by reaction with an acid to form the compound of formula



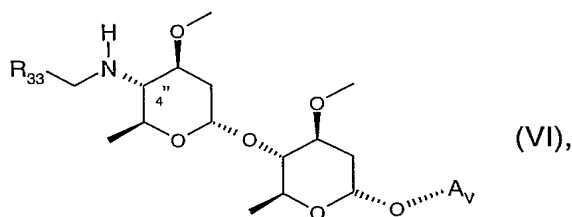
wherein R_1 is as defined hereinbefore;

(E) for the preparation of a compound of formula (I) wherein R_2 is H and R_3 is $-\text{CH}_2-\text{R}_{33}$, with R_{33} being unsubstituted or mono- to penta-substituted C_1-C_{11} alkyl, unsubstituted or mono- to penta-substituted C_2-C_{11} alkenyl; or unsubstituted or mono- to penta-substituted C_2-C_{11} alkynyl; a compound of formula (IV) is reacted first

(E1) with a phosphine or a phosphite; then

(E2) with an aldehyde of formula $\text{R}_{33}-\text{CHO}$; and then

(E3) with a hydride, optionally in the presence of a catalytic amount of acid, to form a compound of formula



wherein A_v corresponds to the macrocyclic structure defined for formula (IV) and R_{33} is as defined hereinbefore; and

(F) the protecting group is split off from the resulting compound of formula (VI) analogously to process step (D); or,

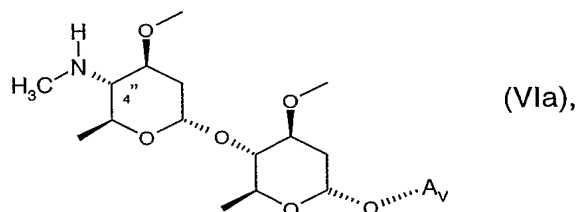
(G) for the preparation of a compound of formula (I) wherein R_2 and R_3 are as defined under process step (E), a compound of formula (V) is reacted analogously to process steps (E1) to (E3);

(H) for the preparation of a compound of formula (I) wherein R_2 is H and R_3 is $-CH_3$, a compound of formula (IV) is reacted first

(H1) with a phosphine; then

(H2) with aqueous formaldehyde in the presence a water-withdrawing agent such as a of a molecular sieve, or with formaldehyde generated from paraformaldehyde; and then

(H3) with a hydride in the presence of a catalytic amount of acid; and the resulting compound of formula



wherein A_v corresponds to the macrocyclic structure defined for formula (IV), is further reacted analogously to process step (D); or,

(I) for the preparation of a compound of formula (I) wherein R_2 is H and R_3 is $-CH_3$, a compound of formula (V) is reacted analogously to process steps (H1) to (H3); or,

(K) for the preparation of a compound of formula (I) wherein R_2 is CH_3 and R_3 is as defined hereinbefore for formula (I), a compound of formula (I) wherein R_3 is as defined hereinbefore for formula (I) is reacted, in accordance with process steps (H2) and (H3) above, first in the presence of a catalytic amount of acid with formaldehyde or paraformaldehyde, and then with a hydride; or,

(L) for the preparation of a compound of formula (I) wherein R_2 is H or unsubstituted or mono- to penta-substituted C_1 - C_{12} alkyl or unsubstituted or mono- to penta-substituted

C₂-C₁₂alkenyl, and R₃ is as defined hereinbefore for formula (I), a compound of formula (I), wherein R₂ is H and R₃ is H or is as defined hereinbefore for formula (I), is reacted with a compound of formula Hal-R₃, wherein R₃ is unsubstituted or mono- to penta-substituted C₁-C₁₂alkyl or unsubstituted or mono- to penta-substituted C₂-C₁₂alkenyl and Hal is halogen, preferably iodine; or,

(M) for the preparation of a compound of formula (I) wherein R₂ and R₃ are CH₃, a compound of formula (I), wherein R₂ and R₃ are hydrogen, is reacted

(M1) with formaldehyde or paraformaldehyde; and then

(M2) with a hydride in the presence of a catalytic amount of acid; or,

(N) for the preparation of a compound of formula (I) wherein R₂ and R₃ together are a three- to seven-membered alkylene or four- to seven-membered alkenylene bridge in each of which a CH₂ group may have been replaced by O, S or NR₄, a compound of formula (I), wherein R₂ and R₃ are H, is reacted with a compound of formula

Hal-C₃-C₇alkylene-Hal or a compound Hal-C₄-C₇alkenylene-Hal,

wherein in the alkylene and alkenylene groups a CH₂ group may have been replaced by O, S or NR₄ and Hal is a halogen.

The invention relates also to

(O) a process for the preparation of a compound of formula (I) wherein R₂ and R₃ are H, in which process either a compound of formula (IV) is reacted, analogously to process step (H1), with a phosphine and then with a base; and the resulting compound, which carries a protecting group in the 5-position, is further reacted analogously to process step (D); or

a compound of formula (V) is reacted, analogously to process step (H1), with a phosphine and then with a base.

The remarks made above in relation to tautomers of compounds of formula (I) apply analogously to starting materials referred to hereinbefore and hereinafter as regards the tautomers thereof.

The reactions described hereinbefore and hereinafter are carried out in a manner known *per se*, for example in the absence or, customarily, in the presence of a suitable solvent or diluent or a mixture thereof, the reactions being carried out, as required, with cooling, at room temperature or with heating, for example in a temperature range of from about -80°C to the boiling point of the reaction medium, preferably from about 0°C to about

+150°C, and, if necessary, in a closed vessel, under pressure, in an inert gas atmosphere and/or under anhydrous conditions. Especially advantageous reaction conditions can be found in the Examples.

The reaction time is not critical; preference is given to a reaction time of from about 0.1 to about 72 hours, especially from about 0.5 to about 24 hours.

The product is isolated by customary methods, for example by means of filtration, crystallisation, distillation or chromatography, or any suitable combination of such methods.

The starting materials mentioned hereinbefore and hereinafter that are used for the preparation of compounds of formula (I) and, where appropriate, tautomers thereof, are known or can be prepared by methods known *per se*, for example in accordance with the information given hereinafter.

Process variant (A):

Examples of suitable solvents and diluents include: aromatic, aliphatic and alicyclic hydrocarbons and halogenated hydrocarbons, such as benzene, toluene, xylene, mesitylene, Tetralin, chlorobenzene, dichlorobenzene, bromobenzene, petroleum ether, hexane, cyclohexane, dichloromethane, trichloromethane, tetrachloromethane, dichloroethane, trichloroethene and tetrachloroethene; ethers, such as diethyl ether, dipropyl ether, diisopropyl ether, dibutyl ether, tert-butyl methyl ether, ethylene glycol monomethyl ether, ethylene glycol monoethyl ether, ethylene glycol dimethyl ether, dimethoxydiethyl ether, tetrahydrofuran and dioxane; or mixtures of the mentioned solvents. Preference is given to dichloromethane.

Suitable leaving groups Q in the compounds of formula (III) are especially sulfonic acid radicals; preference is given, for example, to the anions of toluenesulfonic acid, methanesulfonic acid, trifluoromethanesulfonic acid, pentafluoroethanesulfonic acid and nonafluorobutanesulfonic acid.

Suitable protecting groups Sa in the compounds of formulae (II), (III), (IV) and (VI) are especially trialkylsilyl- or allyloxycarbonyl-groups; preference is given, for example, to trimethylsilyl, triethylsilyl, dimethyl-tert-butylsilyl, dimethyl-isopropylsilyl, dimethyl-1,1,2-trimethylpropylsilyl, diethyl-isopropylsilyl, dimethyl-tert-hexylsilyl; but also to phenyl-tert-alkylsilyl groups such as diphenyl-tert-butylsilyl.

The reactions are advantageously carried out in a temperature range of from about -70°C to +10°C, preferably from -35°C to 0°C.

Especially preferred conditions for the reaction are described in Example A.1.

Process variant (B):

Examples of solvents and diluents include: nitriles, such as acetonitrile; dimethyl sulfoxide; and also alcohols, such as, for example, ethanol or methanol; amides, such as dimethylformamide or dimethylacetamide, are especially suitable.

Especially suitable azide salts are NaN_3 and $\text{Zn}(\text{N}_3)_2 \cdot \text{pyridine}$; especially NaN_3 .

The reactions are advantageously carried out in a temperature range of from -10°C to $+10^\circ\text{C}$.

Especially preferred conditions for the reaction are described, for example, in Example A.1.

Process variant (C):

Examples of suitable solvents and diluents are the same as those mentioned in process variant (A). Also suitable are amides, such as dimethylformamide, dimethylacetamide or hexamethylphosphoric triamide.

Suitable azodicarboxylic acid derivatives are especially azodicarboxylic acid esters, for example the dibenzyl, diethyl, dibutyl, diisopropyl or di-tert-butyl ester or the di-(2,2,2-trichloroethyl) ester; or azodicarboxylic acid amides, such as, for example, N,N,N,N'-azodicarboxylic acid tetramethylamide or azodicarboxylic acid dimorpholide.

Suitable azide sources are especially $(\text{PhO})_2\text{PN}_3$, $\text{Zn}(\text{N}_3)_2 \cdot \text{pyridine}$ or HN_3 .

Suitable phosphines are especially trialkyl- and triaryl-phosphines, such as, for example, trimethyl-, triethyl- and tri-n-butyl-phosphine and also triphenylphosphine.

The reactions are advantageously carried out in a temperature range of from -20°C to 150°C .

Process variant (D):

Examples of suitable solvents and diluents are the same as those mentioned in process variant (A). Also suitable are nitriles, such as acetonitrile; dimethyl sulfoxide; and alcohols, such as, for example, ethanol or methanol; and also water.

The reactions are advantageously carried out in a temperature range of from -10°C to 25°C .

Suitable acids for splitting off the protecting group are, for example, HF in pyridine, $\text{Zn}(\text{BF}_4)_2 \cdot \text{H}_2\text{O}$ or methanesulfonic acid.

Especially preferred conditions for this process variant are described, for example, in Example A.2.

Process variant (E):

Examples of suitable solvents and diluents are the same as those mentioned in process variant (A). Also suitable are nitriles, such as acetonitrile; and also esters of carboxylic acids, such as, for example ethyl acetate.

The reactions are advantageously carried out in a temperature range of from 0°C to +100°C.

Suitable phosphines include, *inter alia*, the same as those mentioned under process variant (C). Suitable phosphites are, for example, trimethyl phosphite, triethyl phosphite, tri-n-butyl phosphite and tri-tert-butyl phosphite.

Suitable hydrides are, especially, complex hydrides, especially sodium borohydride and sodium cyanoborohydride.

Suitable acids are, especially, weak carboxylic acids, such as acetic acid, propionic acid or pivalic acid, especially pivalic acid. The acids are used preferably in catalytic amounts, especially in amounts of less than 10 mol %, more especially less than 5 mol %, very especially less than 2 mol %.

Especially preferred conditions for this process variant are described, for example, in Example A.3.

Process variant (F):

The same process conditions apply as described in variant (D). The reaction is preferably carried out in the presence of methanesulfonic acid in methanol at 0°C.

Process variant (G):

The same process conditions apply as described in variant (E).

Process variant (H):

The same process conditions apply as described in variant (E). As phosphine there are used, for example, trimethylphosphine or tributylphosphine, preferably trimethylphosphine.

In variant (H2), the generation of formaldehyde from paraformaldehyde is carried by heating a suspension of paraformaldehyde in a solvent such as an ether in the presence of a catalytical amount of an acid such as para-toluenesulfonic acid. The formaldehyde is being distilled off the solution.

Process variant (I):

The same process conditions apply as described in variant (H).

Process variant (K):

Preferred solvents are alcohols, such as methanol or ethanol.

The reactions are advantageously carried out in a temperature range of from 0°C to +20°C; preferably from 0°C to 10°C.

Otherwise the same conditions apply as mentioned in process variant (H). Special preference is given to proceeding with sodium borohydride or sodium cyanoborohydride and with pivalic acid as catalyst.

Process variant (L):

The same solvents are used as mentioned in process variant (A). Also suitable are amides, such as dimethylformamide and dimethylacetamide; nitriles, such as acetonitrile; and esters, such as ethyl acetate.

Suitable bases are especially carbonates, such as sodium carbonate, sodium hydrogen carbonate and potassium carbonate, trialkylamines, such as triethylamine, and heterocyclic bases, such as, for example, pyridine.

Process variant (M):

Suitable solvents are especially alcohols, such as methanol or ethanol.

Formaldehyde is preferably used either in the form of an aqueous solution or in the form of paraformaldehyde. Preference is given to a mode of operation in which an excess of formaldehyde or paraformaldehyde is used.

Special preference is given to proceeding with sodium borohydride or sodium cyanoborohydride and with pivalic acid or acetic acid as catalyst.

Process variant (N):

Examples of suitable solvents and diluents are the same as those mentioned in process variant (A). Also suitable are esters of carboxylic acids, such as, for example, ethyl acetate.

Suitable bases are especially carbonates, such as sodium carbonate, sodium hydrogen carbonate or potassium carbonate.

Process variant (O):

Examples of suitable solvents and diluents are the same as those mentioned in process variant (A). Also suitable are nitriles, such as acetonitrile; and also esters of carboxylic acids, such as, for example, ethyl acetate.

Suitable phosphines are especially trialkyl- and triaryl-phosphines, such as, for example, trimethyl- and tri-n-butyl phosphine and also triphenylphosphine.

As bases there are especially used sodium hydroxide solution or ammonium hydroxide, especially in highly diluted form, more especially in a concentration of, for example, 0.01N.

The compounds of formula (I) may be in the form of one of the possible isomers or in the form of a mixture thereof, in the form of pure isomers, or in the form of mixtures of isomers, that is to say in the form of a racemic mixture; the invention relates both to the pure isomers and to the racemic mixtures and this is to be understood accordingly hereinbefore and hereinafter, even when stereochemical details are not specifically mentioned in each case.

The racemates can be resolved into the optical antipodes by known methods, for example by recrystallisation from an optically active solvent, by chromatography on chiral adsorbents, for example high-pressure liquid chromatography (HPLC) on acetyl cellulose, with the aid of suitable microorganisms, by cleavage with specific immobilised enzymes, or *via* the formation of inclusion compounds, for example using chiral crown ethers, in which case only one enantiomer is complexed.

According to the invention, pure optical isomers can be obtained not only by separation of corresponding mixtures of isomers but also by generally known methods of enantio-selective synthesis, for example by carrying out the process according to the invention with starting materials that have appropriate stereochemistry.

It is advantageous to isolate or synthesise whichever isomer is biologically more active, insofar as the individual components have different biological activity.

The compounds of formula (I) may also be obtained in the form of their hydrates and/or may include other solvents, for example solvents which may optionally have been used for the crystallisation of compounds in solid form.

The invention relates to all those embodiments of the process according to which a compound obtainable as starting material or intermediate at any stage of the process is used as starting material and all or some of the remaining steps are carried out, or in which a starting material is used in the form of a derivative and/or a salt and/or its racemates or antipodes, or, especially, is formed under the reaction conditions.

In the process of the present invention it is preferable to use those starting materials and intermediates which result in the compounds of formula (I) that are especially preferred.

The invention relates especially to the preparation processes described in the Examples.

The invention relates also to the compounds of formulae (IV), (V) and (VI), and, where appropriate, E/Z isomers, mixtures of E/Z isomers and/or tautomers, in each case in free form or in salt form.

In the area of pest control, the compounds of formula (I) according to the invention are active ingredients exhibiting valuable preventive and/or curative activity with a very advantageous biocidal spectrum and a very wide spectrum, even at low rates of concentration, while being well tolerated by warm-blooded organisms, fish and plants. The compounds are, surprisingly, equally suitable for controlling plant pests and also ecto- and endo-parasites of humans and especially of productive livestock, domestic animals and pets. They are effective against all or individual development stages of normally sensitive animal pests, but also of resistant animal pests, such as insects and representatives of the order Acarina, nematodes, cestodes and trematodes, whilst at the same time sparing useful organisms. The insecticidal or acaricidal activity of the active ingredients according to the invention may manifest itself directly, i.e. in the mortality of the pests, which occurs immediately or only after some time, for example during moulting, or indirectly, for example in reduced oviposition and/or hatching rate, good activity corresponding to a mortality of at least 50 to 60 %.

The action of the compounds according to the invention and of compositions comprising them against animal pests can be significantly broadened and adapted to prevailing circumstances by the addition of other insecticides, acaricides or nematicides. Examples of suitable additives include representatives of the following classes of active ingredients: organophosphorus compounds, nitrophenols and derivatives, formamidines, ureas, carbamates, pyrethroids, chlorinated hydrocarbons, neonicotinoids and *Bacillus thuringiensis* preparations.

Especially suitable mixing partners are, for example: azamethiphos; chlorfenvinphos; cypermethrin, cypermethrin high-cis; cyromazine; diafenthiuron; diazinon; dichlorvos; dicrotophos; dicyclanil; fenoxycarb; fluazuron; furathiocarb; isazofos; jodfenphos; kinoprene; lufenuron; methacriphos; methidathion; monocrotophos; phosphamidon; profenofos; diofenolan; a compound obtainable from the *Bacillus thuringiensis* strain GC91 or from the strain NCTC11821; pymetrozine; bromopropylate; methoprene; disulfoton; quinalphos; tau-fluvalinate; thiocyclam; thiometon; aldicarb; azinphos-methyl; benfuracarb; bifenthrin; buprofezin; carbofuran; dibutylaminothio; cartap; chlorfluazuron; chlorpyrifos; cyfluthrin; lambda-cyhalothrin; alpha-cypermethrin; zeta-cypermethrin; deltamethrin; diflubenzuron; endosulfan; ethiofencarb; fenitrothion; fenobucarb; fenvalerate; formothion; methiocarb; heptenophos; imidacloprid; isoprocarb; methamidophos; methomyl; mevinphos; parathion; parathion-methyl; phosalone; pirimicarb; propoxur; teflubenzuron; terbufos; triazamate; fenobucarb; tebufenozide; fipronil; beta-cyfluthrin; silafluofen; fenpyroximate; pyridaben; fenazaquin; pyriproxyfen; pyrimidifen; nitenpyram; acetamiprid; avermectin B1 (abamectin); emamectin; emamectin benzoate; spinosad; a plant extract which is active against insects; a preparation comprising nematodes which is active against insects; a preparation obtainable from *Bacillus subtilis*; a preparation comprising fungi which is active against insects; a preparation comprising viruses which is active against insects; chlorfenapyr; acephate; acrinathrin; alanycarb; alphamethrin; amitraz; Az 60541; azinphos A; azinphos M; azocyclotin; bendiocarb; bensultap; beta-cyfluthrin; BPMC; brofenprox; bromophos A; bufencarb; butocarboxim; butylpyridaben; cadusafos; carbaryl; carbophenothion; chloethocarb; chlorethoxyfos; chlormephos; cis-resmethrin; clocythrin; clofentezine; cyanophos; cycloprothrin; cyhexatin; demeton M; demeton S; demeton-S-methyl; dichlofenthion; dicliphos; diethion; dimethoate; dimethylvinphos; dioxathion; edifenphos; esfenvalerate; ethion; ethofenprox; ethoprophos; etrimphos; fenamiphos; fenbutatin oxide; fenothiocarb; fenpropathrin; fenpyrad; fenthion; fluazinam; flucycloxuron; flucythrinate; flufenoxuron; flufenprox; fonophos; fosthiazate; fubfenprox; HCH; hexaflumuron;

hexythiazox; IKI-220; iprobenfos; isofenphos; isoxathion; ivermectin; malathion; mecarbam; mesulfenphos; metaldehyde; metolcarb; milbemectin; moxidectin; naled; Nc 184; omethoate; oxamyl; oxydemeton M; oxydeprofos; permethrin; phenthoate; phorate; phosmet; phoxim; pirimiphos M; pirimiphos E; promecarb; propaphos; prothiofos; prothoate; pyrachlophos; pyridaphenthion; pyresmethrin; pyrethrum; tebufenozide; salithion; sebufos; sulfotep; sulprofos; tebufenpyrad; tebupirimphos; tefluthrin; temephos; terbam; tetrachlorvinphos; thiacloprid; thiafenox; thiodicarb; thiofanox; thionazin; thuringiensin; tralomethrin; triarthene; triazophos; triazuron; trichlorfon; triflumuron; trimethacarb; vamidothion; xylylcarb; YI 5301/5302; zetamethrin; DPX-MP062 – indoxacarb; methoxyfenozide; bifenazate; XMC (3,5-xylyl methylcarbamate); or the fungus pathogen *Metarhizium Anisopliae*; very especially fipronil, thiamethoxam, or lambda-cyhalothrin.

Examples of the abovementioned animal pests which can be controlled with the instantly claimed compounds are:

from the order Lepidoptera, for example,

Acleris spp., *Adoxophyes* spp., *Aegeria* spp., *Agrotis* spp., *Alabama argillaceae*, *Amylois* spp., *Anticarsia gemmatilis*, *Archips* spp., *Argyrotaenia* spp., *Autographa* spp., *Busseola fusca*, *Cadra cautella*, *Carposina nipponensis*, *Chilo* spp., *Choristoneura* spp., *Clysia ambiguella*, *Cnaphalocrocis* spp., *Cnephasia* spp., *Cochylis* spp., *Coleophora* spp., *Crocidolomia binotalis*, *Cryptophlebia leucotreta*, *Cydia* spp., *Diatraea* spp., *Diparopsis castanea*, *Earias* spp., *Ephestia* spp., *Eucosma* spp., *Eupoecilia ambiguella*, *Euproctis* spp., *Euxoa* spp., *Grapholita* spp., *Hedya nubiferana*, *Heliothis* spp., *Hellula undalis*, *Hyphantria cunea*, *Keiferia lycopersicella*, *Leucoptera scitella*, *Lithocollethis* spp., *Lobesia botrana*, *Lymantria* spp., *Lyonetia* spp., *Malacosoma* spp., *Mamestra brassicae*, *Manduca sexta*, *Operophtera* spp., *Ostrinia nubilalis*, *Pammene* spp., *Pandemis* spp., *Panolis flammea*, *Pectinophora gossypiella*, *Phthorimaea operculella*, *Pieris rapae*, *Pieris* spp., *Plutella xylostella*, *Prays* spp., *Scirpophaga* spp., *Sesamia* spp., *Sparganothis* spp., *Spodoptera* spp., *Synanthedon* spp., *Thaumetopoea* spp., *Tortrix* spp., *Trichoplusia ni* and *Yponomeuta* spp.;

from the order Coleoptera, for example,

Agriotes spp., *Anthonomus* spp., *Atomaria linearis*, *Chaetocnema tibialis*, *Cosmopolites* spp., *Curculio* spp., *Dermestes* spp., *Diabrotica* spp., *Epilachna* spp., *Eremnus* spp., *Leptinotarsa decemlineata*, *Lissorhoptrus* spp., *Melolontha* spp., *Oryzaephilus* spp., *Otiorhynchus* spp., *Phlyctinus* spp., *Popillia* spp., *Psylliodes* spp., *Rhizopertha* spp., *Scarabeidae*, *Sitophilus* spp., *Sitotroga* spp., *Tenebrio* spp., *Tribolium* spp. and *Trogoderma* spp.;

from the order Orthoptera, for example,

Blatta spp., *Blattella* spp., *Gryllotalpa* spp., *Leucophaea maderae*, *Locusta* spp., *Periplaneta* spp. and *Schistocerca* spp.;

from the order Isoptera, for example, *Reticulitermes* spp.;

from the order Psocoptera, for example, *Liposcelis* spp.;

from the order Anoplura, for example, *Haematopinus* spp., *Linognathus* spp., *Pediculus* spp., *Pemphigus* spp. and *Phylloxera* spp.;

from the order Mallophaga, for example, *Damalinea* spp. and *Trichodectes* spp.;

from the order Thysanoptera, for example, *Frankliniella* spp., *Hercinothrips* spp., *Taeniothrips* spp., *Thrips palmi*, *Thrips tabaci* and *Scirtothrips aurantii*;

from the order Heteroptera, for example, *Cimex* spp., *Distantiella theobroma*, *Dysdercus* spp., *Euchistus* spp., *Eurygaster* spp., *Leptocorisa* spp., *Nezara* spp., *Piesma* spp., *Rhodnius* spp., *Sahlbergella singularis*, *Scotinophara* spp. and *Triatoma* spp.;

from the order Homoptera, for example, *Aleurothrixus floccosus*, *Aleyrodes brassicae*, *Aonidiella* spp., *Aphididae*, *Aphis* spp., *Aspidiotus* spp., *Bemisia tabaci*, *Ceroplaster* spp., *Chrysomphalus aonidium*, *Chrysomphalus dictyospermi*, *Coccus hesperidum*, *Empoasca* spp., *Eriosoma larigerum*, *Erythroneura* spp., *Gascardia* spp., *Laodelphax* spp., *Lecanium corni*, *Lepidosaphes* spp., *Macrosiphus* spp., *Myzus* spp., *Nephotettix* spp., *Nilaparvata* spp., *Paratoria* spp., *Pemphigus* spp., *Planococcus* spp., *Pseudaulacaspis* spp., *Pseudococcus* spp., *Psylla* spp., *Pulvinaria aethiopica*, *Quadraspidotus* spp., *Rhopalosiphum* spp., *Saissetia* spp., *Scaphoideus* spp., *Schizaphis* spp., *Sitobion* spp., *Trialeurodes vaporariorum*, *Trioza erytrae* and *Unaspis citri*;

from the order Hymenoptera, for example, *Acromyrmex*, *Atta* spp., *Cephus* spp., *Diprion* spp., *Diprionidae*, *Gilpinia polytoma*, *Hoplocampa* spp., *Lasius* spp., *Monomorium pharaonis*, *Neodiprion* spp., *Solenopsis* spp. and *Vespa* spp.;

from the order Diptera, for example, *Aedes* spp., *Antherigona soccata*, *Bibio hortulanus*, *Calliphora erythrocephala*, *Ceratitis* spp., *Chrysomyia* spp., *Culex* spp., *Cuterebra* spp., *Dacus* spp., *Drosophila melanogaster*, *Fannia* spp., *Gastrophilus* spp., *Glossina* spp., *Hypoderma* spp., *Hyppobosca* spp., *Liriomyza* spp., *Lucilia* spp., *Melanagromyza* spp., *Musca* spp., *Oestrus* spp., *Orseolia* spp., *Oscinella frit*, *Pegomyia hyoscyami*, *Phorbia* spp., *Rhagoletis pomonella*, *Sciara* spp., *Stomoxys* spp., *Tabanus* spp., *Tannia* spp. and *Tipula*

spp.;

from the order Siphonaptera, for example, *Ctenocephalides felis*, *Ctenocephalides canis*, *Ceratophyllus* spp. and *Xenopsylla cheopis*;

from the order Thysanura, for example, *Lepisma saccharina* and

from the order Acarina, for example, *Acarus siro*, *Aceria sheldoni*, *Aculus schlechtendali*, *Amblyomma* spp., *Argas* spp., *Boophilus* spp., *Brevipalpus* spp., *Bryobia praetiosa*, *Calipitimerus* spp., *Chorioptes* spp., *Dermanyssus gallinae*, *Eotetranychus carpini*, *Eriophyes* spp., *Hyalomma* spp., *Ixodes* spp., *Olygonychus pratensis*, *Ornithodoros* spp., *Panonychus* spp., *Phyllocoptura oleivora*, *Polyphagotarsonemus latus*, *Psoroptes* spp., *Rhipicephalus* spp., *Rhizoglyphus* spp., *Sarcoptes* spp., *Tarsonemus* spp. and *Tetranychus* spp.;

from the class Trematoda especially representatives of the family Fasciolidae, especially *Fasciola hepatica*;

from the class Nematoda, for example,

root knot nematodes, cyst-forming nematodes and also stem and leaf nematodes; especially of *Heterodera* spp., e.g. *Heterodera schachtii*, *Heterodera avenae* and *Heterodera trifolii*; *Globodera* spp., e.g. *Globodera rostochiensis*; *Meloidogyne* spp., e.g. *Meloidogyne incognita* and *Meloidogyne javanica*; *Radopholus* spp., e.g. *Radopholus similis*; *Pratylenchus*, e.g. *Pratylenchus neglectans* and *Pratylenchus penetrans*; *Tylenchulus*, e.g. *Tylenchulus semipenetrans*; *Longidorus*, *Trichodorus*, *Xiphinema*, *Ditylenchus*, *Aphelenchoides* and *Anguina*; especially *Meloidogyne*, e.g. *Meloidogyne incognita*, and *Heterodera*, e.g. *Heterodera glycines*.

An especially important aspect of the present invention is the use of the compounds of formula (I) according to the invention for the protection of plants against parasitic feeding pests.

The active ingredients according to the invention can be used to control, i.e. to inhibit or destroy, pests of the mentioned type occurring especially on plants, more especially on useful plants and ornamentals in agriculture, in horticulture and in forestry, or on parts of such plants, such as the fruits, blossoms, leaves, stems, tubers or roots, while in some cases parts of plants that grow later are still protected against those pests.

Target crops are especially cereals, e.g. wheat, barley, rye, oats, rice, maize and sorghum; beet, such as sugar beet and fodder beet; fruit, e.g. pomes, stone fruit and soft

fruit, such as apples, pears, plums, peaches, almonds, cherries and berries, e.g. strawberries, raspberries and blackberries; leguminous plants, such as beans, lentils, peas and soybeans; oil plants, such as rape, mustard, poppy, olives, sunflowers, coconut, castor oil, cocoa and groundnuts; cucurbitaceae, such as marrows, cucumbers and melons; fibre plants, such as cotton, flax, hemp and jute; citrus fruits, such as oranges, lemons, grapefruit and mandarins; vegetables, such as spinach, lettuce, asparagus, cabbages, carrots, onions, tomatoes, potatoes and paprika; lauraceae, such as avocado, cinnamon and camphor; and tobacco, nuts, coffee, aubergines, sugar cane, tea, pepper, vines, hops, bananas, natural rubber plants and ornamentals.

Further areas of use of the active ingredients according to the invention are the protection of stored goods and storerooms and the protection of raw materials, and also in the hygiene sector, especially the protection of domestic animals and productive livestock against pests of the mentioned type, especially against infestation of domestic animals, especially cats and dogs, by fleas, ticks and nematodes.

The invention therefore also relates to a pesticidal composition which comprises a compound of the formula (I), in the form of emulsifiable concentrates, suspension concentrates, directly sprayable or dilutable solutions, spreadable pastes, dilute emulsions, wettable powders, soluble powders, dispersible powders, wettable powders, dusts, granules or encapsulations in polymeric substances, all of which comprise - at least - one of the active ingredients according to the invention and which are to be selected to suit the intended aims and prevailing circumstances.

In these compositions, the active ingredient is employed together with - at least - one of the auxiliaries conventionally used in the art of formulation, such as extenders, e.g. solvents or solid carriers, or such as surface-active compounds (surfactants).

Examples of suitable solvents are: non-hydrogenated or partially hydrogenated aromatic hydrocarbons, preferably the fractions C₈ to C₁₂ of alkylbenzenes, such as xylene mixtures, alkylated naphthalenes or tetrahydronaphthalene, aliphatic or cycloaliphatic hydrocarbons such as paraffins or cyclohexane, alcohols such as ethanol, propanol or butanol, glycols and their ethers and esters such as propylene glycol, dipropylene glycol ether, ethylene glycol, ethylene glycol monomethyl ether or ethylene glycol monoethyl ether, ketones such as cyclohexanone, isophorone or diacetone alcohol, strongly polar solvents such as N-methylpyrrolid-2-one, dimethyl sulfoxide or N,N-dimethylformamide, water, free or epoxidized rapeseed, castor, coconut or soya oil, and silicone oils.

Solid carriers which are used for example for dusts and dispersible powders are, as a rule, ground natural minerals, such as calcite, talc, kaolin, montmorillonite or attapulgite. To improve the physical properties, it is also possible to add highly-disperse silicas or highly-disperse absorptive polymers. Suitable particulate adsorptive carriers for granules are porous types, such as pumice, brick grit, sepiolite or bentonite, and suitable non-sorptive carrier materials are calcite or sand. Moreover, a large number of granulated materials of inorganic or organic nature can be used, in particular dolomite or comminuted plant residues.

Suitable surface-active compounds are, depending on the nature of the active ingredient to be formulated, non-ionic, cationic and/or anionic surfactants or surfactant mixtures which have good emulsifying, dispersing and wetting properties. The surfactants listed below are only to be considered as examples; many more surfactants conventionally used in the art of formulation and suitable in accordance with the invention are described in the relevant literature.

Suitable non-ionic surfactants are primarily polyglycol ether derivatives of aliphatic or cycloaliphatic alcohols, of saturated or unsaturated fatty acids and alkylphenols which can contain 3 to 30 glycol ether groups and 8 to 20 carbon atoms in the (aliphatic) hydrocarbon radical and 6 to 18 carbon atoms in the alkyl radical of the alkylphenols. Also suitable are water-soluble polyethylene oxide adducts with polypropylene glycol, ethylenediamino-polypropylene glycol and alkylpolypropylene glycol having 1 to 10 carbons in the alkyl chain and 20 to 250 ethylene glycol ether and 10 to 100 propylene glycol ether groups. The abovementioned compounds normally contain 1 to 5 ethylene glycol units per propylene glycol unit. Examples which may be mentioned are nonylphenylpolyethoxyethanols, castor oil polyglycol ethers, polypropylene/polyethylene oxide adducts, tributylphenoxypolyethoxyethanol, polyethylene glycol and octylphenoxypolyethoxyethanol. Also suitable are fatty acid esters of polyoxyethylene sorbitan, such as polyoxyethylene sorbitan trioleate.

The cationic surfactants are mainly quaternary ammonium salts which have, as substituents, at least one alkyl radical of 8 to 22 carbon atoms and, as further substituents, lower alkyl, benzyl or lower hydroxyalkyl radicals which may be halogenated. The salts are preferably in the form of halides, methylsulfates or ethylsulfates. Examples are stearyltrimethylammonium chloride and benzyldi(2-chloroethyl)ethylammonium bromide.

Suitable anionic surfactants can be both water-soluble soaps and water-soluble synthetic surface-active compounds. Soaps which are suitable are the alkali metal salts,

alkaline earth metal salts and unsubstituted or substituted ammonium salts of higher fatty acids (C_{10} - C_{22}), such as the sodium or potassium salts of oleic or stearic acid, or of natural fatty acid mixtures which can be obtained, for example, from coconut or tall oil; mention must also be made of the fatty acid methyltaurates. However, synthetic surfactants, in particular fatty sulfonates, fatty sulfates, sulfonated benzimidazole derivatives or alkylarylsulfonates, are used more frequently. As a rule, the fatty sulfonates and fatty sulfates exist as alkali metal salts, alkaline earth metal salts or unsubstituted or substituted ammonium salts and generally have an alkyl radical of 8 to 22 C atoms, alkyl also including the alkyl moiety of acyl radicals; examples which may be mentioned are the sodium or calcium salt of lignosulfonic acid, of the dodecylsulfuric ester or of a fatty alcohol sulfate mixture prepared with natural fatty acids. This group also includes the salts of the sulfuric esters and sulfonic acids of fatty alcohol/ethylene oxide adducts. The sulfonated benzimidazole derivatives preferably contain 2 sulfo groups and one fatty acid radical having approximately 8 to 22 C atoms. Examples of alkylarylsulfonates are the sodium, calcium or triethanolammonium salts of dodecylbenzenesulfonic acid, of dibutyl-naphthalenesulfonic acid or of a naphthalenesulfonic acid/formaldehyde condensate. Also suitable are corresponding phosphates, such as salts of the phosphoric ester of a p-nonylphenol (4-14)ethylene oxide adduct, or phospholipids.

As a rule, the compositions comprise 0.01 to 99.99%, in particular 0.1 to 99%, in particular 0.1 to 95%, of active ingredient and 0.01 to 99.99%, especially 1 to 99.9%, in particular 5 to 99.9%, of - at least - one solid or liquid auxiliary, it being possible, as a rule, for the surfactants to amount to 0 to 25%, in particular 0.1 to 20%, of the composition (% is in each case per cent by weight). While concentrated compositions are more preferred as commercially available goods, the end user uses, as a rule, dilute compositions, which have a considerably lower concentration of active ingredient. Preferred compositions are composed in particular as follows (% = per cent by weight):

Emulsifiable concentrates:

active ingredient:	1 to 90%, preferably 5 to 20%
surfactant:	1 to 30%, preferably 10 to 20 %
solvent:	5 to 98%, preferably 70 to 85%

Dusts:

active ingredient:	0.1 to 10%, preferably 0.1 to 1%
solid carrier:	99.9 to 90%, preferably 99.9 to 99%

Suspension concentrates:

active ingredient:	5 to 75%, preferably 10 to 50%
water:	94 to 24%, preferably 88 to 30%
surfactant:	1 to 40%, preferably 2 to 30%

Wettable powder:

active ingredient:	0.5 to 90%, preferably 1 to 80%
surfactant:	0.5 to 20%, preferably 1 to 15%
solid carrier:	5 to 99%, preferably 15 to 98%

Granules:

active ingredient:	0.5 to 30%, preferably 3 to 15%
solid carrier:	99.5 to 70%, preferably 97 to 85%

The activity of the compositions according to the invention can be widened considerably, and adapted to prevailing circumstances, by adding other insecticidally, acaricidally and/or fungicidally active ingredients. Examples of suitable additives of active ingredients are representatives of the following classes of active ingredients: organophosphorus compounds, nitrophenols and derivatives, formamidines, ureas, carbamates, pyrethroids, chlorinated hydrocarbons and *Bacillus thuringiensis* preparations. The compositions according to the invention may also comprise other solid or liquid auxiliaries such as stabilizers, for example epoxidized or unepoxidized vegetable oils (e.g. epoxidized coconut oil, rapeseed oil or soya oil), antifoams, e.g. silicone oil, preservatives, viscosity regulators, binders and/or tackifiers, and also fertilizers or other active ingredients for achieving specific effects, for example bactericides, nematocides, molluscicides or selective herbicides.

The compositions according to the invention are prepared in a known manner, in the absence of auxiliaries for example by grinding and/or screening a solid active ingredient, or active ingredient mixture, for example to obtain a specific particle size, and in the presence of at least one auxiliary for example by intimately mixing and/or grinding the active ingredient, or active ingredient mixture, with the auxiliary(or auxiliaries). These processes for the preparation of the compositions according to the invention and the use of the compounds of the formula (I) for the preparation of these compositions are also subjects of the invention.

The methods of application for the compositions, i.e. the methods of controlling pests of the abovementioned type, such as spraying, atomizing, dusting, brushing on, seed-dressing, scattering or pouring, which are to be selected to suit the intended aims and the prevailing circumstances, and the use of the compositions for controlling pests of the abovementioned type are also subjects of the invention. Typical use concentrations are between 0.1 and 1000 ppm, preferably between 0.1 and 500 ppm, of active ingredient. The rates of application per hectare are generally 1 to 2000 g of active ingredient per hectare, in particular 10 to 1000 g/ha, preferably 20 to 600 g/ha, especially 20 to 100 g/ha.

A preferred method of application in the field of crop protection is application to the foliage of the plant (foliar application), it being possible to adjust frequency and rate of application to the degree or risk of infestation with the pest in question. Alternatively, the active ingredient can reach the plants via the root system (systemic action), by drenching the locus of the plants with a liquid composition or by incorporating the active ingredient in solid form into the locus of the plants, for example in the soil, for example in the form of granules (soil application). In the case of paddy rice crops, such granules can be metered into the flooded paddy field.

The compositions according to the invention are also suitable for the protection of plant propagation material, e.g. seed such as fruits, tubers or kernels, or nursery plants, against fungal infection and animal pests. The propagation material can be dressed with the composition before planting, for example seed can be dressed before sowing. It is also possible to apply the active ingredients according to the invention to seed kernels (coating), either by soaking the kernels in a liquid composition or by coating them with a solid composition. Alternatively, the composition can be applied to the site of application when the propagation material is planted, for example into the seed furrow during sowing. These treatment methods for plant propagation material and the plant propagation material treated accordingly are further subjects of the invention.

Preparation Examples

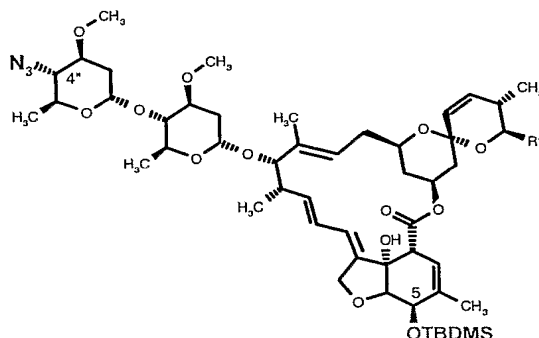
In the following Examples, the preparation of avermectin B1 derivatives (mixtures of avermectin B1a and B1b derivatives) is described. The B1b derivative generally is present in the mixtures only in an amount of about from 5 to 10 % by weight, for which reason only the bands of the B1a derivative can usually be seen in the NMR spectrum.

The abbreviations in the details of NMR data denote:

s: singlet, MHz: megahertz, brs: broad singlet; t: triplet; m: multiplet; d: doublet;
J: coupling constant.

TBDMS in the Examples denotes the radical $-\text{Si}(\text{CH}_3)_2(\text{tert-butyl})$, R_1 denotes a mixture of isopropyl and sec-butyl.

Example A.1): Preparation of 4"-deoxy-4"-(S)-azido-5-OTBDMS-avermectin B1 of formula

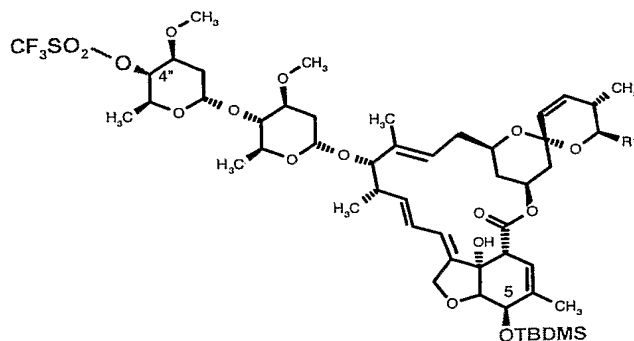


Synthesis variant a):

To a solution of 10.93 g of 4''-(*R*)-5-OTBDMS-avermectin B1 (mixture of B1a and B1b derivative) in 180 ml of absolute tetrahydrofuran under argon there are added 11.61 g of triphenylphosphine and the mixture is cooled to 0°C. 13.21 g of dibenzyl azodicarboxylate and 12.19 g of diphenylphosphoryl azide in 120 ml of absolute tetrahydrofuran are added, dropwise, to the reaction solution over the course of 30 minutes, with stirring. The mixture is allowed to warm up to room temperature and is then heated at 50°C for 5 hours. The mixture is left to stand overnight at room temperature and is then poured onto ice-water and extracted three times with ether. Washing twice with saturated aqueous NaCl solution and drying over Na₂SO₄ and concentration by evaporation yields the crude product in the form of a viscous oil. Column chromatography on silica gel using ethyl acetate/hexane (1:10 to 1:4) as eluant, dissolving in CH₂Cl₂ and subsequent evaporation of the solvent and drying under a high vacuum yields the desired product in the form of a colourless dry foam.

Synthesis variant b):

Step 1): To 987 mg of 4''-(*R*)-5-OTBDMS-avermectin B1 in 30 ml of absolute CH₂Cl₂ under argon there are added, in rapid succession, at -35°C, 732 mg of 4-N,N-dimethylaminopyridine, 775 mg of ethyldiisopropylamine (Hünig's base) and 1129 mg of trifluoromethanesulphonic anhydride, with vigorous stirring, and stirring is continued at 0°C for 2 h. After 70 minutes, a further portion of 0.2 ml of trifluoromethanesulphonic anhydride is added. The mixture is then poured onto ice-water, extracted twice with ether, washed several times each with water and with saturated aqueous NaCl solution and dried with Na₂SO₄, and the solvent is evaporated off. Fractionated filtration over silica gel on a column using ethyl acetate/hexane (1:2) as eluant yields the compound of formula



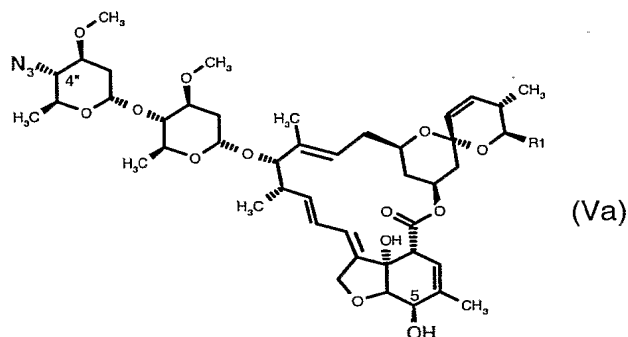
in the form of a pale-yellow foam.

Step 2): To 883 mg of the resulting intermediate in 5 ml of absolute dimethylformamide there are added, with ice-H₂O-cooling, 102 mg of sodium azide and stirring is carried out for a further 15 hours. Aqueous working-up using ethyl acetate, water and saturated aqueous

NaCl solution, drying (Na_2SO_4) and column-chromatographic purification of the crude product on silica gel using ethyl acetate/hexane (1:5) as eluant yields 493 mg of 4"-deoxy-4"-(S)-azido-5-OTBDMS-avermectin B1b and 61 mg of a mixture of 4"-deoxy-4"-(S)-azido-5-OTBDMS-avermectin B1a+1b, both in the form of a colourless dry foam.

In NMR (500 MHz), the title product exhibits the following selected signals (δ , ppm) for the B1a derivative: 0.13 (6H, s, $\text{Si}(\text{CH}_3)_2$), 1.49 (3H, brs, CH_3 -14a), 1.78 (3H, brs, CH_3 -4a), 2.98 (1H, t, $J = 9.7\text{Hz}$, CH-4"), 3.20 (1H, t, $J = 9.0\text{Hz}$, CH-4'), 3.39 (1H, m, CH-2), 3.42 (3H, s, OCH_3), 3.46 (3H, s, OCH_3), 3.82 (d, $J = 5.7\text{Hz}$, CH-6), 3.92 (1H, brs, CH-13), 4.09 (1H, s, HO-C-7), 4.43 (1H, m, CH-5), 4.56-4.69 (2H, ABX system, $J_1 = 1.5\text{Hz}$, $J_2 = 14.4\text{Hz}$, CH_2 -8a), 4.75 (1H, d, $J = 3.0\text{Hz}$, CH-1'), 4.99 (1H, m, CH-15), 5.32 (1H, d, $J = 1\text{Hz}$, CH-3), 5.40 (1H, d, $J = 3.0\text{Hz}$, CH-1").

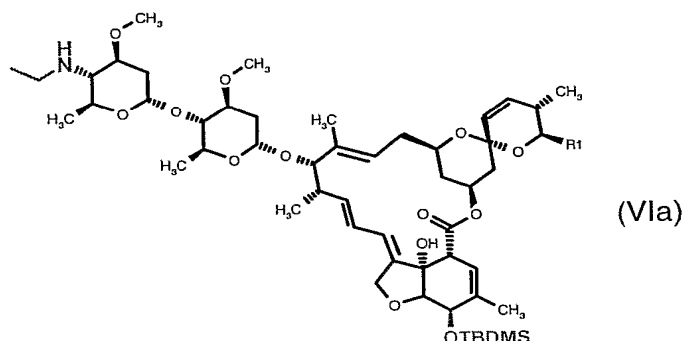
Example A.2): Preparation of 4"-deoxy-4"-(S)-azido-avermectin B1 of formula



To a solution of 805 mg of 4"-deoxy-4"-(S)-azido-5-OTBDMS-avermectin B1 of formula (IVa) in 5 ml of absolute THF under argon there are added, with ice-cooling, 20 ml of HF-pyridine reagent in tetrahydrofuran, and the mixture is left to stand at room temperature for 22 hours. The reaction mixture is poured onto cooled saturated aqueous NaHCO_3 solution and extracted twice with ether. The combined organic extracts are washed with water and with saturated aqueous NaCl solution (twice in each case), dried over MgSO_4 and concentrated by evaporation. The crude product is purified by column chromatography on silica gel using ethyl acetate/hexane (2:5) as eluant.

The title product is thus obtained, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.48 (3H, brs, CH_3 -14a), 1.87 (3H, brs, CH_3 -4a), 2.98 (1H, t, $J = 9.8\text{Hz}$, CH-4"), 3.20 (1H, t, $J = 9.0\text{Hz}$, CH-4'), 3.29 (1H, m, CH-2), 3.42 (3H, s, OCH_3), 3.46 (3H, s, OCH_3), 3.92 (1H, brs, CH-13), 3.96 (1H, d, $J = 6.1\text{Hz}$, CH-6), 4.29 (1H, brd, $J = 5.9\text{Hz}$, CH-5), 4.64-4.72 (2H, ABX system, CH_2 -8a), 4.76 (1H, d, $J = 3\text{Hz}$, CH-1'), 4.98 (1H, dm, $J = 7.3\text{Hz}$, CH-15).

Example A.3): Preparation of 4"-deoxy-4"-(S)-ethylamino-5-OTBDMS-avermectin B1 of formula



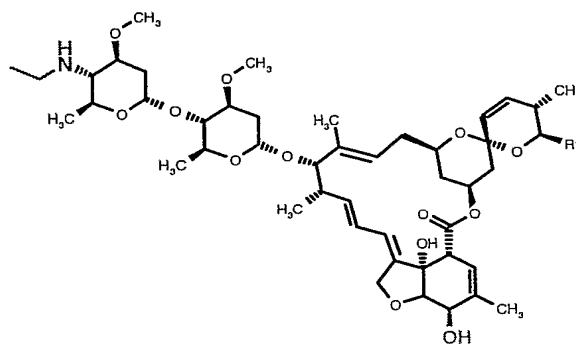
Step 1): 1 g of 4"-deoxy-4"-(S)-azido-5-OTBDMS-avermectin B1 in 50 ml of absolute tetrahydrofuran under argon is stirred together with 2.97 ml of trimethylphosphine (1M in tetrahydrofuran) for 4 hours at 65°C.

Step 2): To the resulting solution from Step 1) there is added 0.224 ml of acetaldehyde and the mixture is stirred at 40°C for 15 hours. The solution is then concentrated by evaporation *in vacuo*.

Step 3): The crude product from the previous Step is taken up in 30 ml of methanol; 45 mg of NaBH₄ are added and the mixture is stirred at room temperature for 2 hours. The solution is poured onto water and extracted with ethyl acetate. Subsequent washing of the organic phase with saturated aqueous NaCl solution and drying with Na₂SO₄ yields the crude product after concentration by evaporation. Purification is carried out on silica gel using CH₂Cl₂/MeOH (95:5) as eluant.

4"-Deoxy-4"-(S)-ethylamino-5-OTBDMS-avermectin B1 is thus obtained, which, in NMR (500 MHz), exhibits the following signals (δ , ppm) for the B1a derivative: 0.12 (6H, s, 5-OSi(CH₃)₂), 1.09 (3H, t, $J = 7\text{Hz}$, CH₃CH₂NH), 1.49 (brs, CH₃-14a), 1.78 (3H, brs, CH₃-4a), 2.17 (t, $J = 9.5\text{Hz}$, CH-4"), 2.69 (2H, m, CH₃CH₂NH), 3.22 (1H, t, $J = 9\text{Hz}$, CH-4'), 3.38 (3H, s, OCH₃), 3.42 (3H, s, OCH₃), 3.82 (1H, d, $J = 5.7\text{Hz}$, CH-6), 3.92 (1H, brs, CH-13), 4.43 (1H, m, CH-5), 4.56-4.69 (2H, ABX system, CH₂-8a), 4.99 (1H, m, CH-15).

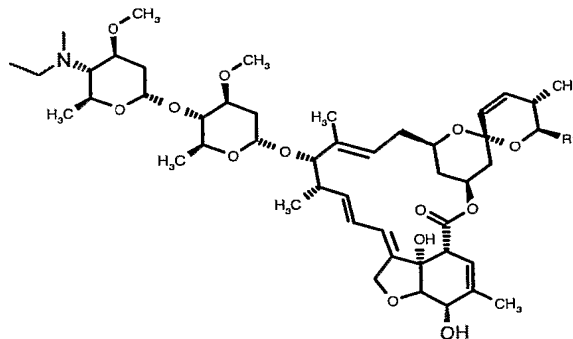
Example A.4): 4"-Deoxy-4"-(S)-ethylamino-avermectin B1 of formula



484 mg of 4"-deoxy-4"-(S)-ethylamino-5-OTBDMS-avermectin B1 in 30 ml of methanol under argon are treated for 2 hours with 60 mg of methanesulfonic acid with ice/water cooling. The reaction solution is added to saturated NaHCO_3 solution, extracted three times with ethyl acetate, washed with saturated aqueous NaCl solution and dried over Na_2SO_4 . The crude product, concentrated by evaporation, is purified on silica gel using ethyl acetate/hexane (3:1) as eluant.

Drying *in vacuo* yields 4"-deoxy-4"-(S)-ethylamino-avermectin B1, which, in NMR (500 MHz), exhibits the following signals (δ , ppm) for the B1a derivative: 1.09 (3H, t, $J = 7.1\text{ Hz}$, $\text{CH}_3\text{CH}_2\text{NH}$), 1.48 (brs, CH_3 -14a), 1.86 (3H, brs, CH_3 -4a), 2.16 (t, $J = 9.4\text{ Hz}$, CH-4"), 2.68 (2H, m, $\text{CH}_3\text{CH}_2\text{NH}$), 3.22 (1H, t, $J = 8.9\text{ Hz}$, CH-4'), 3.29 (1H, m, CH-2), 3.38 (3H, s, OCH_3), 3.43 (3H, s, OCH_3), 3.92 (brs, CH-13), 3.96 (d, $J = 6.4\text{ Hz}$, CH-6), 4.28 (1H, m, CH-5), 4.58-4.76 (2H, ABX system, CH_2 -8a), 4.75 (1H, d, $J = \text{approx. } 3\text{ Hz}$, CH-1'), 4.98 (1H, m, CH-15).

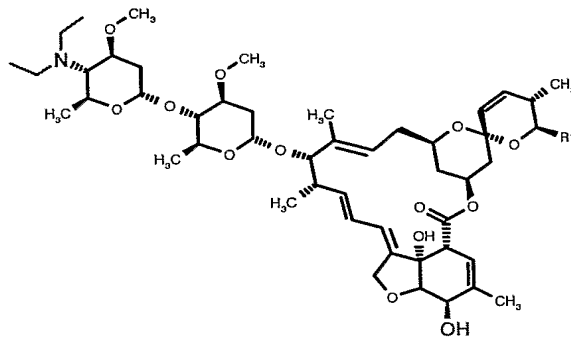
Example A.5): Preparation of 4"-deoxy-4"-(S)-N-ethyl-N-methyl-avermectin B1 of formula



To a solution of 130 mg of 4"-deoxy-4"-(S)-ethylamino-avermectin B1 in 4 ml of methanol under argon there are added, with stirring, a catalytic amount of pivalic acid and 0.310 ml of formaldehyde (37 % in water). After 1.75 hours, 11 mg of NaCNBH₃ are added and stirring is carried out for a further 2 hours. The reaction solution is poured onto saturated aqueous NaHCO₃ solution, extracted three times with ethyl acetate, washed three times with saturated aqueous NaCl solution and dried over Na₂SO₄. The crude product, concentrated by evaporation, is purified on silica gel using ethyl acetate/hexane (1:2) as eluant and dried *in vacuo*.

There is thus obtained 4"-deoxy-4"-(S)-N-ethyl-N-methyl-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.02 (3H, t, $J = 7.1\text{Hz}$, $\text{CH}_3\text{CH}_2\text{N}$) 1.48 (3H, brs, CH₃-14a), 1.87 (3H, brs, CH₃-4a), 2.21 (t, $J = 9.7\text{Hz}$, CH-4"), 2.33 (3H, s, NCH₃), 2.65 (2H, q, $J = 7.1\text{Hz}$, $\text{CH}_3\text{CH}_2\text{N}$), 3.24 (1H, t, $J = 9\text{Hz}$), 3.29 (1H, m, CH-2), 3.35 (3H, s, OCH₃), 3.43 (3H, s, OCH₃), 3.93 (brs, CH-13), 3.97 (d, $J = 6.1\text{Hz}$), 4.01 (1H, s, HO-C-7), 4.28 (1H, m, CH-5), 4.64-4.71 (2H, ABX system, CH₂-8a), 4.76 (1H, d, $J = 4.2\text{Hz}$, CH'-1'), 4.99 (1H, m, CH-15).

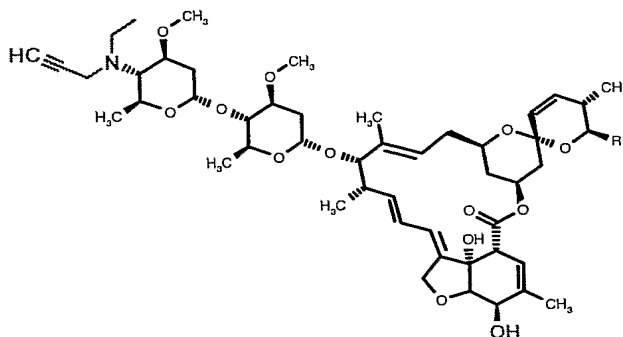
Example A.6): Preparation of 4"-deoxy-4"-(S)-N,N-diethyl-avermectin B1 of formula



To a two-phase mixture of 50 mg of 4"-deoxy-4"-(S)-ethylamino-avermectin B1 in 5 ml of ethyl acetate and 5.6 ml of 0.1M NaHCO₃ in H₂O there is added 0.56 ml of ethyl iodide. The reaction mixture is stirred vigorously in a closed vessel at 60-68°C for 72 hours; it is then poured onto water, extracted with ethyl acetate, washed with saturated aqueous NaCl solution and subsequently dried over Na₂SO₄. The crude product, concentrated by evaporation, is separated from the unreacted starting material on silica gel using ethyl acetate/hexane (1:1) as eluant.

There is thus obtained 4"-deoxy-4"-(S)-N,N-diethylamino-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.1 (6H, t, $J = 7\text{Hz}$, $(\text{CH}_3\text{CH}_2)_2\text{N}$), 1.48 (3H, brs, CH_3 -14a), 1.86 (3H, brs, CH_3 -4a), 2.31 (t, $J = \text{approx. } 10\text{Hz}$, CH-4"), 2.67 (4H, m, $(\text{CH}_3\text{CH}_2)_2\text{N}$), 3.26 (1H, t, $J = 9\text{Hz}$, CH-4'), 3.31 (1H, m, CH-2), 3.36 (3H, s, OCH_3), 3.44 (3H, s, OCH_3), 4.30 (1H, m, CH-5), 4.66-4.74 (2H, ABX system, CH_2 -8a).

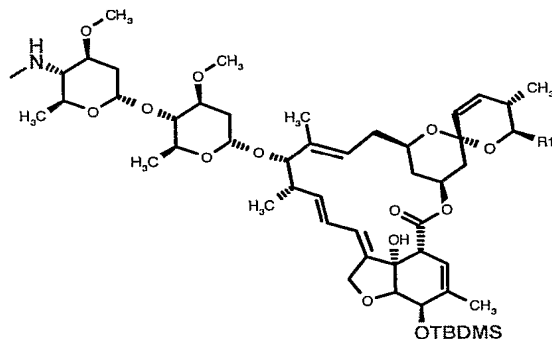
Example A.7): Preparation of 4"-deoxy-4"-(S)-N-ethyl-N-propargylamino-avermectin B1 of formula



To a two-phase mixture of 50 mg of 4"-deoxy-4"-(S)-ethylamino-avermectin B1 in 3 ml of ethyl acetate and 5.6 ml of 0.1M NaHCO_3 in H_2O there is added 0.42 ml of propargyl bromide. The reaction mixture is stirred vigorously in a closed vessel at 70°C for 72 hours; it is then poured onto water, extracted with ethyl acetate, washed with saturated aqueous NaCl solution and then dried over Na_2SO_4 . The crude product, concentrated by evaporation, is purified on silica gel using ethyl acetate/hexane (1:1) as eluant.

There is thus obtained the title product 4"-deoxy-4"-(S)-N-ethyl-N-propargylamino-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.06 (3H, m, $\text{CH}_3\text{CH}_2\text{N}$), 1.49 (brs, CH_3 -14a), 1.87 (3H, brs, CH_3 -4a), 2.15 (1H, brs, $\text{CH}\equiv\text{CHCH}_2\text{N}$), 2.28 (t, $J = 10\text{Hz}$, CH-4"), 2.33 (d, $J = 8\text{Hz}$, HO-C-5), 2.65-2.85 (2H, dm, $\text{CH}_3\text{CH}_2\text{N}$), 3.23 (1H, t, $J = 9\text{Hz}$, CH-4'), 3.29 (1H, m, CH-2), 3.33 (3H, s, OCH_3), 3.43 (3H, s, OCH_3), 3.49 (brs, $\text{CH}\equiv\text{CHCH}_2\text{N}$), 3.93 (brs, CH-13), 3.97 (d, $J = 6.4\text{Hz}$, CH-6), 4.01 (1H, s, HO-C-7), 4.28 (1H, m, CH-5), 4.64-4.71 (2H, ABX system, CH_2 -8a), 5.00 (1H, m, CH-15).

Example A.8): Preparation of 4"-deoxy-4"-(S)-methylamino-5-OTBDMS-avermectin B1 of formula



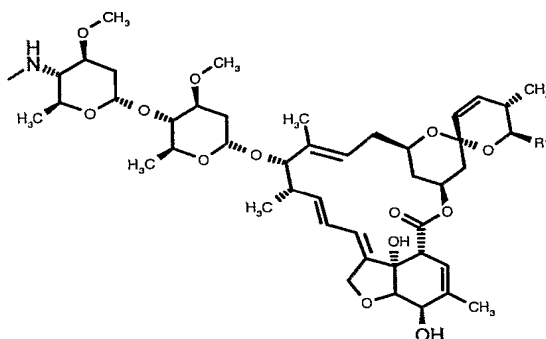
Step 1): A solution of 500 mg of 4"-deoxy-4"-(S)-azido-5-OTBDMS-avermectin B1 and 1.48 ml of trimethylphosphine (1M in tetrahydrofuran) in 40 ml of absolute tetrahydrofuran under argon is left to stand at room temperature for 72 hours.

Step 2): To the resulting solution there are added approximately 15 g of 0.4 nm molecular sieve (in bead form) and also 0.320 ml of formaldehyde (37 % in H₂O), and the mixture is stirred at 65°C for 24 hours. Filtration is carried out and the solution is concentrated by evaporation *in vacuo* and dried to constant weight. The imine intermediate is obtained in the form of a colourless dry foam.

Step 3): The intermediate from Step 2) is taken up in 30 ml of methanol; a spatula tip of pivalic acid and 22.5 mg of NaBH₄ are added and stirring is carried out at room temperature for 2 hours. The mixture is then poured onto water, extracted with ethyl acetate, washed with saturated NaCl solution and dried over Na₂SO₄. The crude product obtained is purified on silica gel using CH₂Cl₂/MeOH (95:5) as eluant.

There is thus obtained 4"-deoxy-4"-(S)-methylamino-5-OTBDMS-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 0.12 (6H, s, Si(CH₃)₂), 1.49 (brs, CH₃-14a), 1.78 (brs, CH₃-4a), 2.10 (1H, t, J = 9.5Hz, CH-4"), 2.46 (3H, s, CH₃NH), 3.23 (t, J = 9Hz, CH-4'), 3.38 (3H, s, OCH₃), 3.42 (3H, s, OCH₃), 3.92 (1H, brs, CH-13), 4.42 (1H, m, CH-5), 4.56-4.69 (2H, ABX system, J_1 = 2.2Hz, J_2 = 14.2Hz, CH₂-8a), 4.76 (1H, d, J = 3.9Hz, CH-1'), 4.99 (1H, m, CH-15).

Example A.9): Preparation of 4"-deoxy-4"-(S)-methylamino-avermectin B1 of formula



Synthesis variant a)

A solution of 250 mg of 4"-deoxy-4"-(S)-methylamino-5-OTBDMS-avermectin B1 in 20 ml of methanol is treated, at 0-5°C, with 31 mg of methanesulfonic acid for 5 hours. The reaction solution is added to saturated NaHCO₃ solution. The mixture is extracted three times with ethyl acetate, washed three times with saturated NaCl solution and dried with Na₂SO₄. 230 mg of crude product are obtained. Purification on silica gel using CH₂Cl₂/MeOH (95:5) as eluant yields the desired product in the form of a colourless dry foam after drying under a high vacuum.

Synthesis variant b)

Step 1): 300 mg of 4"-deoxy-4"-(S)-azido-avermectin B1 are stirred in 25 ml of absolute tetrahydrofuran under argon with 1.67 ml of trimethylphosphine (1M in tetrahydrofuran) for 12 hours at room temperature and then for 24 hours at 65°C.

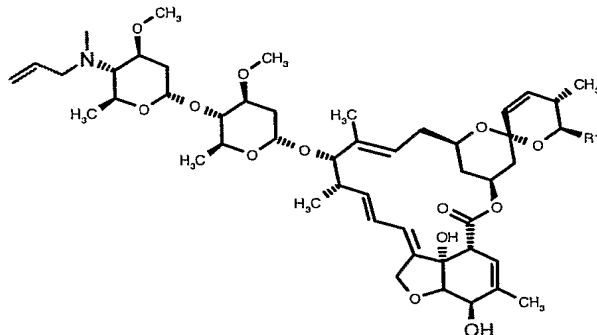
Step 2): To the resulting solution there are added 15 g of 0.4 nm molecular sieve (in bead form) and also 0.217 ml of formaldehyde (37 % in H₂O) and the mixture is stirred at 70°C for 24 hours.

Step 3): The intermediate from Step 2) is dissolved in 15 ml of methanol, and 15 mg of NaBH₄ are added. After stirring for 5 hours, the mixture is poured onto water, extracted with ethyl acetate, washed with saturated aqueous NaCl solution and dried over Na₂SO₄. The crude product is purified on silica gel using CH₂Cl₂/MeOH (95:5) as eluant.

There is thus obtained 4"-deoxy-4"-(S)-methylamino-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.15 (3H, d, $J = 7.1$ Hz), 1.23 (3H, d, $J = 6.1$ Hz), 1.27 (3H, d, $J = 6.4$ Hz), 1.48 (3H, brs, CH₃-14a), 1.86 (3H, brs, CH₃-4a), 2.09 (1H, t, $J = 9.6$ Hz, CH-4"), 2.46 (3H, s, CH₃NH), 3.23 (1H, t, $J = 9$ Hz,

CH-4'), 3.29 (1H, m, CH-2), 3.38 (3H, s, OCH₃), 3.43 (3H, s, OCH₃), 3.92 (1H, brs, CH-13), 3.96 (1H, d, $J = 6.2\text{Hz}$, CH-6), 4.29 (1H, dm, $J = 6.2\text{ Hz}$, CH-5), 4.64 – 4.71 (2H, ABX system, CH₂-8a), 4.99 (1H, m, CH-15).

Example A.10): Preparation of 4"-deoxy-4"-(S)-N-2-propenyl-N-methylamino-avermectin B1 of formula

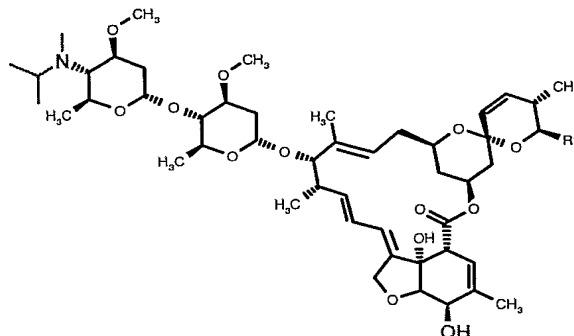


To a two-phase solution consisting of 84 mg of 4"-deoxy-4"-(S)-methylamino-avermectin B1 in 5 ml of ethyl acetate and 9.5 ml of 0.1M NaHCO₃ in H₂O there is added 0.80 ml of allyl bromide. The reaction mixture is stirred vigorously in a closed vessel, first for 18 hours at room temperature and then for 12 hours at 40°C. The mixture is then poured onto water, extracted with ethyl acetate, washed with saturated aqueous NaCl solution and dried over Na₂SO₄. The crude product, concentrated by evaporation, is purified on silica gel using ethyl acetate/hexane (1:1) as eluant.

There is thus obtained 4"-deoxy-4"-(S)-N-2-propenyl-N-methylamino-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.48 (brs, CH₃-14a), 1.86 (3H, brs, CH₃-4a), 2.23 (t, $J = 9.7\text{Hz}$, CH-4"), 2.30 (s, NCH₃), 3.24 (t, $J = 9.1\text{Hz}$, CH-4'), 3.29 (m, CH-2), overlapped by 3.28 (m, NCH₂CH=CH₂), 3.36 (3H, s, OCH₃), 3.43 (3H, s, OCH₃), 3.93 (brs, CH-13), 3.96 (1H, d, $J = 6.3\text{Hz}$, CH-6), 4.01 (1H, s, HO-C(7)), 4.28 (1H, m, CH-5), 4.64-4.70 (2H, ABX system, CH₂-8a), 4.75 (1H, d, $J = \text{approx. } 3\text{Hz}$, CH-1'), 4.99 (1H, m, CH-15), 5.03-5.16 (2H, m, NCH₂CH=CH₂).

Example P:11): Preparation of 4"-deoxy-4"-(S)-N-isopropyl-N-methylamino-avermectin

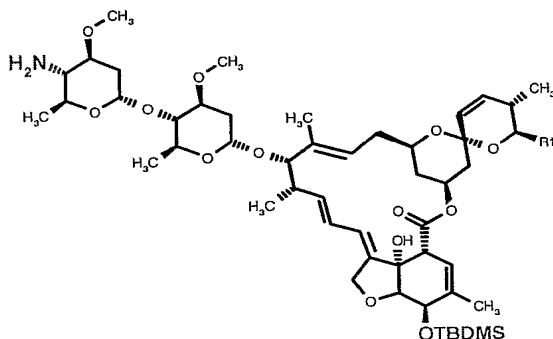
B1 of formula



To a mixture consisting of 100 mg of 4"-deoxy-4"-(S)-methylamino-avermectin B1 in 5 ml of ethyl acetate and 11.3 ml of NaHCO₃ (0.1M in H₂O) there is added 0.226 ml (20 eq.) of 2-iodopropane, and heating is carried out, with vigorous stirring, at 75°C for 72 h. The mixture is poured onto water, extracted with ethyl acetate, washed with saturated NaCl solution and dried with Na₂SO₄. From that crude mixture there is isolated, besides unreacted starting material, the desired compound (column chromatography on silica gel using CH₂Cl₂/MeOH (9:1) as eluant).

There is thus obtained 4"-deoxy-4"-(S)-N-isopropyl-N-methylamino-avermectin B1a, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.03 (6H, t, $J = 6.6$ Hz, NCH(CH₃)₂), 1.48 (brs, CH₃-14a), 1.87 (3H, brs, CH₃-4a), 2.31 (t, $J = 9.7$ Hz, CH-4"), 2.32 (d, $J = 8.3$ Hz, HO-C-5), 2.91 (1H, sep, $J = 6.6$ Hz, NCH(CH₃)₂), 3.25 (1H, t, $J = 8.9$ Hz, CH-4'), 3.29 (1H, m, CH-2), 3.33 (3H, s, OCH₃), 3.43 (3H, s, OCH₃), 3.93 (brs, CH-13), 3.96 (1H, d, 6.1Hz, CH-6), 4.01 (1H, s, HO-C-7), 4.28 (1H, m, CH-5), 4.64-4.71 (2H, ABX system, CH₂-8a), 4.76 (1H, d, $J = \text{approx. } 3$ Hz, CH-1'), 4.99 (1H, m, CH-15).

Example A.12): Preparation of 4"-deoxy-4"-(S)-amino-5-OTBDMS-avermectin B1 of formula

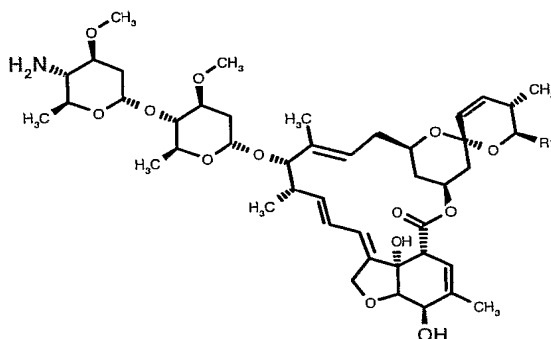


Step 1): A solution of 500 mg of 4"-deoxy-4"-(S)-azido-5-OTBDMS-avermectin B1 and 1.48 ml of trimethylphosphine (1M in tetrahydrofuran) in 40 ml of absolute tetrahydrofuran under argon is left to stand at room temperature for 72 hours.

Step 2): To the solution from Step 1) there are added 6 ml of 0.1N aqueous ammonium hydroxide, and the mixture is left to stand for 15 hours. The solution is poured onto water, extracted three times with ethyl acetate, neutralised with water and a small amount of dilute sulfuric acid, washed with saturated aqueous NaCl solution and dried over Na₂SO₄. The crude product, concentrated by evaporation, is purified on silica gel using CH₂Cl₂/MeOH (95:5) as eluant.

There is thus obtained 4"-deoxy-4"-(S)-amino-5-OTBDMS-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 0.13 (6H, s, Si(CH₃)₂), 1.49 (brs, CH₃-14a), 1.78 (brs, CH₃-4a), 2.46 (t, J = 9.4Hz, CH-4"), 3.22 (1H, t, J = 9Hz, CH-4'), 3.40 (s, OCH₃), 3.43 (s, OCH₃), 3.82 (d, J = 6.6Hz, CH-6), 3.92 (1H, brs, CH-13), 4.05-4.13 (2H, br, NH₂), 4.42 (1H, m, CH-5), 4.56-4.69 (2H, ABX system, CH₂-8a), 4.76 (1H, d, J = approx. 3Hz), 4.99 (1H, m, CH-15).

Example A.13): Preparation of 4"-deoxy-4"-(S)-amino-avermectin B1 of formula



a) Variant 1

255 mg of 4"-deoxy-4"-(S)-amino-5-OTBDMS-avermectin B1 and 32 mg of methanesulfonic acid in 20 ml of methanol are stirred at 0°C for 5 hours. The solution is poured onto water, extracted with ethyl acetate, washed with saturated aqueous NaCl solution and dried (Na₂SO₄). The crude product, concentrated by evaporation, is purified on silica gel using CH₂Cl₂/MeOH (95:5) as eluant. After drying, the title product is obtained.

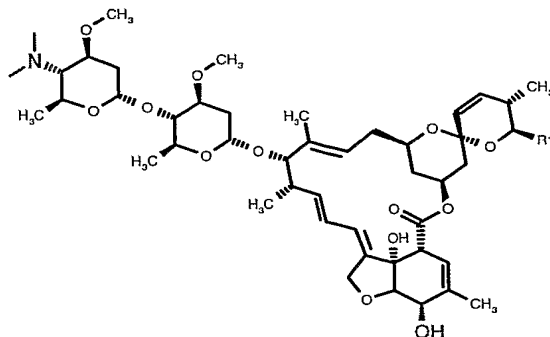
b) Variant 2

Step 1): 1.41 g of 4"-deoxy-4"-(S)-azido-avermectin B1 in 100 ml of absolute tetrahydrofuran under argon are stirred with 7.85 ml of trimethylphosphine (1M in tetrahydrofuran) at 50°C for 48 hours.

Step 2): To the solution from Step 1) there are added 30 ml of NaOH (0.001N) and heating is carried out at 45°C for 18 hours. The mixture is poured onto water and extracted three times with ethyl acetate. The organic phase is washed with saturated aqueous NaCl solution and dried over Na₂SO₄. Purification of the crude product is carried out on silica gel using CH₂Cl₂/MeOH (95:5) as eluant.

There is thus obtained 4"-deoxy-4"-(S)-amino-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.48 (brs, CH₃-14a), 1.86 (brs, CH₃-4a), 2.46 (t, $J = 9.4$ Hz, CH-4"), 3.22 (1H, t, $J = 9$ Hz, CH-4'), 3.29 (1H, m, CH-2), 3.39 (s, OCH₃), 3.43 (s, OCH₃), 3.93 (brs, CH-13), 3.96 (d, $J = 6.1$ Hz, CH-6), 4.28 (1H, m, CH-5), 4.64-4.71 (2H, ABX system, CH₂-8a), 4.76 (1H, d, $J = \text{approx. } 3$ Hz, CH-1'), 4.99 (1H, m, CH-15).

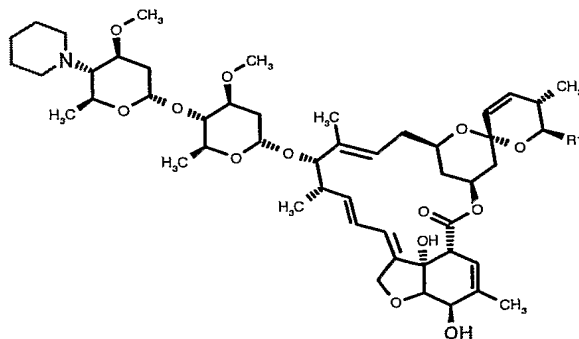
Example A.14): Preparation of 4"-deoxy-4"-(S)-N,N-dimethylamino-avermectin B1 of formula



123 mg of 4"-deoxy-4"-(S)-amino-avermectin B1 and 100 mg of pivalic acid in 2 ml of acetonitrile are treated, for 2 hours, with 57 mg of formaldehyde (37 % in H₂O) at room temperature. The reaction mixture is then poured onto saturated aqueous NaHCO₃ solution and extracted twice with ethyl acetate. The combined organic extracts are washed with saturated aqueous NaCl solution (twice) and dried over Na₂SO₄. The crude product is purified on silica gel in ethyl acetate/hexane (1:3) to (2:1). The title product is obtained in the form of a colourless, dry foam.

There is thus obtained 4"-deoxy-4"-(S)-N,N-dimethylamino-avermectin B1, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.48 (brs, CH₃-14a), 1.87 (3H, brs, CH₃-4a), 2.14 (1H, t, $J = 9.7$ Hz, CH-4"), 2.41 (6H, s, (CH₃)₂N), 3.24 (1H, t, $J = 9$ Hz, CH-4'), 3.29 (1H, m, CH-2), 3.35 (3H, s, OCH₃), 3.43 (3H, s, OCH₃), 3.93 (1H, brs, CH-13), 3.96 (1H, d, $J = 6.4$ Hz, CH-6), 4.01 (1H, s, HO-C-7), 4.28 (1H, m, CH-5), 4.64-4.70 (2H, ABX system, CH₂-8a), 4.75 (1H, d, $J = \text{approx. } 3$ Hz, CH-1'), 4.99 (1H, m, CH-15).

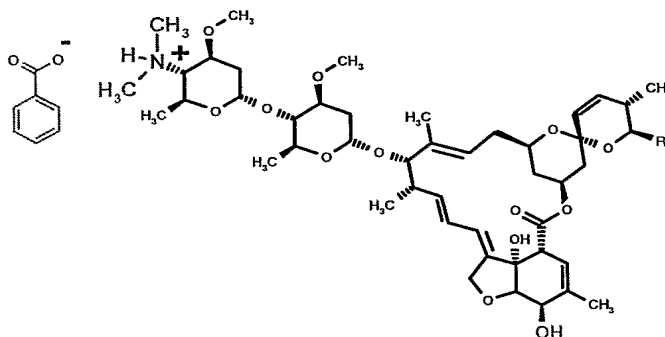
Example A.15): Preparation of 4"-deoxy-4"-(S)- (piperidin-1-yl)-avermectin B1



A solution of 100 mg of 4"-deoxy-4"-(S)-amino-avermectin B1, 40 mg of 1,5-dibromopentane and 0.016 ml of triethylamine in a mixture of 4 ml of THF and 1 ml of ethyl acetate is stirred at room temperature for 3 days. The solution is poured onto water, extracted with ethyl acetate, washed with saturated aqueous NaCl solution and dried over Na₂SO₄. The desired product is isolated from the crude mixture with the aid of column chromatography on silica gel in CH₂Cl₂/MeOH (9:1) as eluant.

There is thus obtained 4"-deoxy-4"-(S)- (piperidin-1-yl)-avermectin, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.48 (brs, CH₃-14a), 1.87 (brs, CH₃-4a), 2.06 (t, J = 9.8Hz, CH-4"), 2.32 (d, J = 8.3Hz, HO-C-5), 2.56 and 2.77 (2 m, 4H, N(CH₂)₂), 3.23 (1H, t, J = 9Hz, CH-4'), 3.36 (3H, s, OCH₃), 3.42 (3H, s, OCH₃), 3.93 (1H, brs, CH-13), 3.96 (1H, d, J = 6.1Hz, CH-6), 4.01 (1H, s, HO-C-7), 4.28 (1H, m, CH-5), 4.64-4.71 (2H, ABX system, CH₂-8a), 4.75 (1H, d, J = approx. 3Hz, CH-1'), 4.99 (1H, m, CH-15).

Example A.16): Preparation of the benzoate salt of 4"-deoxy-4"-(S)-N,N-dimethyl-amino-avermectin B1.



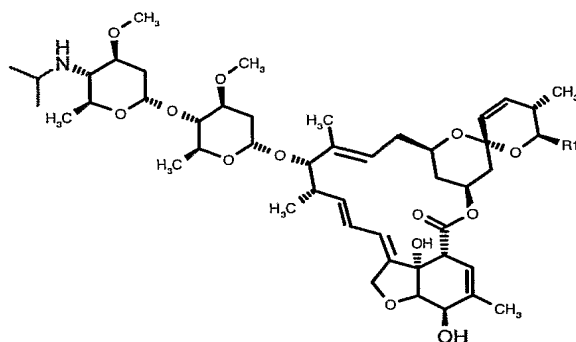
100 mg of 4"-deoxy-4"-(S)-N,N-dimethylamino-avermectin B1 and 14 mg of benzoic acid are dissolved in 20 ml of diethyl ether and concentrated to dryness by evaporation. The desired salt is thus obtained.

In analogous manner, the 1:1 salts of 4"-deoxy-4"-(S)-N,N-dimethylamino-avermectin B1 with isobutyric acid, salicylic acid, 2,5-dimethylbenzoic acid, thiosalicylic acid, 4-phenoxybutyric acid and citric acid are also prepared.

In analogous manner to Example A.16, the 1:1 salts of 4"-deoxy-4"-(S)-N-methylamino-avermectin B1 with benzoic acid, isobutyric acid, salicylic acid, 2,5-

dimethylbenzoic acid, thiosalicylic acid, 4-phenoxybutyric acid and citric acid are also prepared.

Example A.17: Preparation of 4''-deoxy-4''-(S)-N-isopropylamino-avermectin B₁



150 mg of 4''-(S)-amino-avermectin B₁ in 5 ml of THF are kept at room temperature for 15 hours together with 0.050 ml of acetone and 0.3 ml of ethyl acetate/water (9:1). Treatment with 8.5 mg of NaBH₄ is then carried out for 4 hours. The mixture is poured onto an ice-water mixture and extracted 3 times with ethyl acetate. The combined organic extracts are washed with saturated sodium chloride solution and dried with Na₂SO₄. Purification is carried out on silica gel using CH₂Cl₂/MeOH (95:5) as eluant.

There is thus obtained the title product, which, in NMR (500 MHz), exhibits the following selected signals (δ , ppm) for the B1a derivative: 1.05 (d overlapped, CH(CH₃)₂), 1.48 (brs, CH₃-14a), 1.86 (3H, brs, CH₃-4a), 2.24 (t, $J = 10.3\text{Hz}$, CH-4''), 2.90 (1H, m, NCH(CH₃)₂), 3.21 (t, $J = 9.1\text{Hz}$, CH-4'), 3.29 (1H, m, CH-2), 3.38 (3H, s, OCH₃), 3.43 (3H, s, OCH₃), 3.92 (1H, brs, CH-13), 3.96 (1H, d, $J = 6.3\text{Hz}$, CH-6), 3.99 (1H, s, HO-C(7)), 4.29 (1H, m, CH-5), 4.64-4.71 (2H, ABX system, CH₂-8a), 4.76 (1H, d, $J = \text{approx. } 3\text{Hz}$, CH-1'), 4.99 (1H, m, CH-15).

The compounds listed in Tables A and 1 to 6 can also be prepared analogously to the above Preparation Examples. In Tables 1 to 6, the symbol \sim indicates the bond by way of which the corresponding radical is attached to the nitrogen atom of the basic structure.

Because the compounds are, in most cases, present as mixtures of the avermectin derivatives B1a and B1b, characterisation by means of the usual physical data, such as melting point and refractive index, makes little sense. The compounds are therefore characterised, as indicated in the above Examples, after chromatographic purification, by

means of NMR spectroscopy or using the retention times that are determined in HPLC analysis (high-resolution liquid chromatography). The indication B1a in the Tables relates to the main component, wherein R₁ is sec-butyl, the content of which is usually greater than 80 %. B1b stands for the subsidiary component, wherein R₁ is isopropyl. In the case of compounds where a retention time is indicated only in the B1a column, it is not possible to determine the retention time for the B1b component because of the low B1b derivative content. Assignment of the correct structures of the B1a and B1b components is carried out by mass spectrometry.

The following method is used for the HPLC analysis:

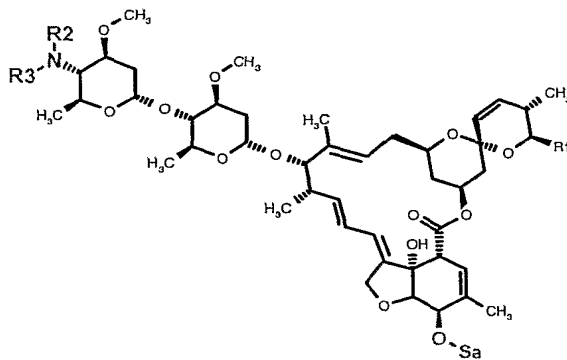
HPLC gradient conditions				
solvent A:	0.01 % trifluoroacetic acid in H ₂ O			
solvent B:	0.01% trifluoroacetic acid in CH ₃ CN			
time [min]	A [%]	B [%]	flow rate 1 [μl/min]	flow rate 2 [ml/min]
0	80	20	500	3,5
0.1	50	50	500	3,5
10	5	95	500	3,5
15	0	100	500	3,5
17	0	100	500	3,5
17.1	80	20	500	3,5
22	80	20	500	3,5

column:	YMC-Pack ODS-AQ 1	YMC-Pack ODS-AQ 2
column length:	125 mm	30 mm
column internal diameter:	2 mm	4.6 mm
temperature:	40°C	40 °C

The YMC-Pack ODS-AQ column used for chromatography of the compounds is manufactured by YMC, Alte Raesfelderstrasse 6, 46514 Schermbeck, Germany. For the HPLC analysis of compounds A.18 to A.34 of table A and of table 1, and for the compounds

1.70 and 1,159, flow rate 1 and YMC-Pack ODS-AQ 1 conditions were used, for all other compounds, flow rate 2 and YMC-Pack ODS-AQ 2 conditions were used.

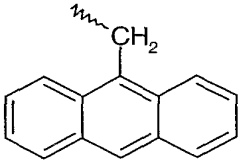
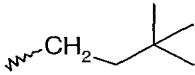
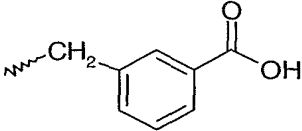
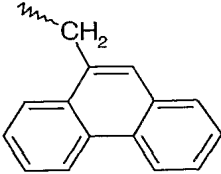
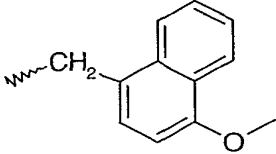
Table A: Compounds of formula

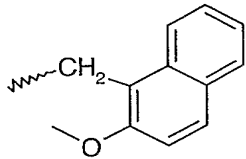
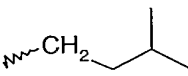
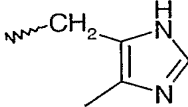
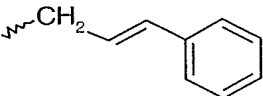
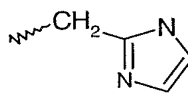
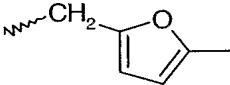
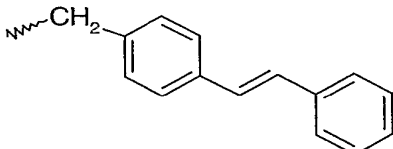


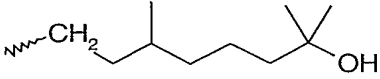
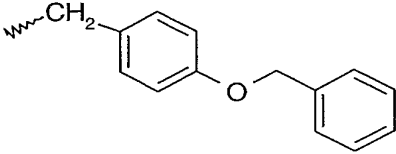
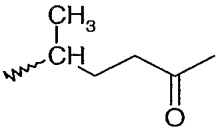
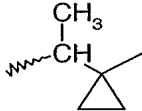
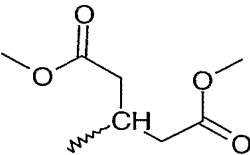
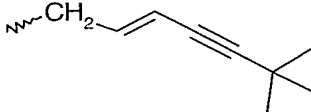
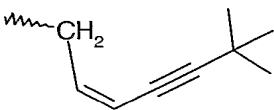
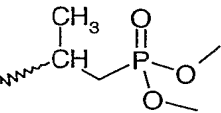
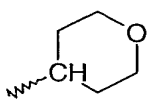
wherein R₁ is sec-butyl (B1a) or isopropyl (B1b).

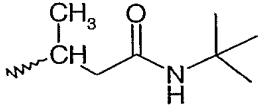
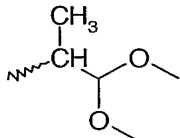
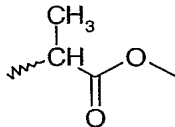
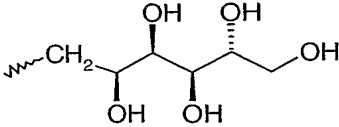
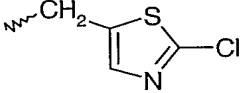
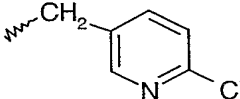
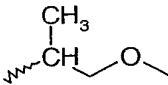
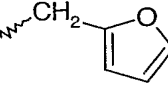
	R ₂	R ₃	Sa	LC B _{1a} (min)	LC B _{1b} (min)
A.18	H	CH ₃ CH ₂ CH ₂	TBDMS	9.87	-
A.19	H	CH ₃ CH ₂ CH ₂	H	5.47	-
A.20	H	PhCH ₂	TBDMS	10.03	-
A.21	H	PhCH ₂	H	5.92	-
A.22	CH ₃	PhCH ₂	H	6.72	6.13
A.23	CH ₃	CH ₃ CH ₂ CH ₂	H	5.57	-
A.24	CH ₃	propargyl	H	6.67	6.14
A.25	CH ₃	C ₂ H ₅ OCOCH ₂	H	7.74	7.15
A.26	CH ₃	CH ₃ OCOCH ₂	H	7.63	7.09
A.27	CH ₃	4-F-PhCH ₂	H	7.10	6.56
A.28	C ₂ H ₅	C ₂ H ₅ OCOCH ₂	H	8.81	-
A.29	C ₂ H ₅	allyl	H	6.00	5.51
A.30	C ₃ H ₇ (n)	propargyl	H	8.94	8.40
A.31	C ₃ H ₇ (n)	C ₂ H ₅	H	5.98	-
A.32	C ₃ H ₇ (n)	C ₂ H ₅ OCOCH ₂	H	9.82	9.34
A.33		-(CH ₂) ₄ -	H	6.88	6.38
A.34		-(CH ₂) ₃ -	H	5.11	4.75

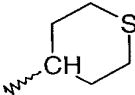
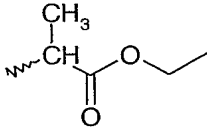
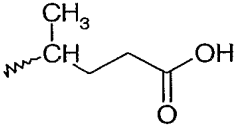
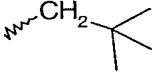
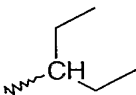
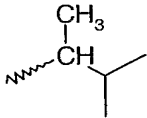
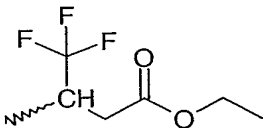
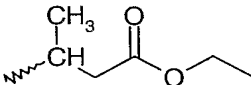
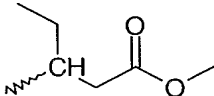
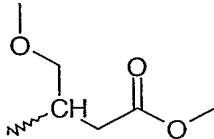
Table 1: Compounds of formula (I) wherein R₁ is sec-butyl (B1a) or isopropyl (B1b) and R₂ is hydrogen.

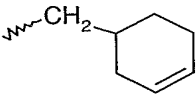
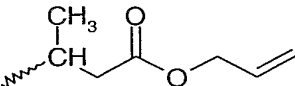
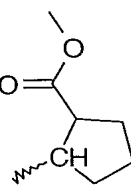
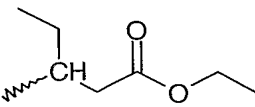
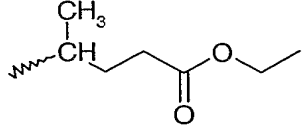
No.	R ₃	Retention time (min.)	
		B1a	B1b
1.001			
1.002	(2-hydroxy-3-methoxy-phenyl)methyl		
1.003	(3-nitrophenyl)methyl		
1.004	(4-dimethylaminophenyl)methyl		
1.005	(3-bromophenyl)methyl		
1.006	(2-nitro-5-hydroxy-phenyl)methyl		
1.007	2-phenyl-n-propyl		
1.008	3-pyridyl-methyl		
1.009	(2,4-dimethylphenyl)methyl		
1.010	(3-fluorophenyl)methyl		
1.011			
1.012			
1.013			
1.014	(3,4-methylenedioxy-6-nitrophenyl)methyl		
1.015			
1.016	(2-hydroxy-5-methoxyphenyl)methyl		
1.017	(2-hydroxy-4-methoxyphenyl)methyl		
1.018	(2,3-dihydroxyphenyl)methyl		
1.019	(2-hydroxy-5-nitrophenyl)methyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.020			
1.021	dec-4-en-1-yl		
1.022	(3-[4-methoxyphenoxy]-phenyl)methyl		
1.023			
1.024	3-phenyl-n-propyl		
1.025	4-pyridyl-methyl		
1.026			
1.027	ethyl		
1.028	n-butyl	2.48	2.32
1.029			
1.030	(2-bromophenyl)methyl		
1.031			
1.032			
1.033	(4-n-propoxyphenyl)methyl		
1.034			
1.035	(2-chloro-4-hydroxyphenyl)methyl		
1.036	cyclododecyl		
1.037	1-methyl-n-butyl		
1.038	4-hydroxy-1-methyl-n-butyl		
1.039	1-methyl-n-propyl		

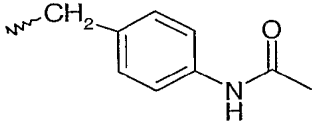
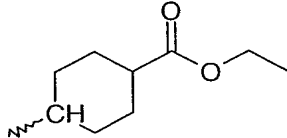
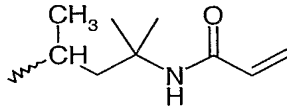
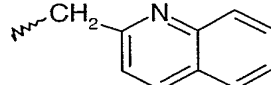
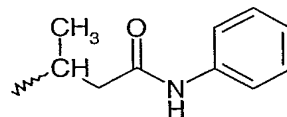
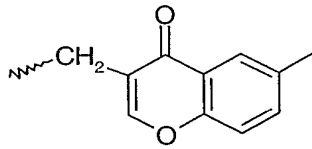
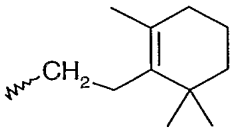
No.	R ₃	Retention time (min.)	
		B1a	B1b
1.040			
1.041	(4-tert-butyl-phenyl)methyl		
1.042	3-phenyl-n-butyl		
1.043			
1.044	-CH ₂ -C(=O)-O-CH ₂ -phenyl	2.70	-
1.045	-CH ₂ -C(=O)-O-methyl	2.40	2.23
1.046	-CH ₂ -C(=O)-O-ethyl	2.44	2.35
1.047			
1.048			
1.049			
1.050			
1.051			
1.052			
1.053			
1.054	1,2-dimethyl-3-hydroxy-n-propyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.055	cyclobutyl		
1.056			
1.057			
1.058	-CH ₂ C(=O)-O-isopropyl		
1.059	-CH ₂ -C(=O)-O-CH ₂ -CH ₂ -O-CH ₃		
1.060	-CH ₂ C(=O)-O-tert-butyl	2.61	2.48
1.061	-CH ₂ C(=O)NH ₂	2.19	-
1.062			
1.063			
1.064	propargyl		
1.065	2-chloroallyl		
1.066	3,3-dichloroallyl		
1.067			
1.068			
1.069			
1.070		5.45	5.07
1.071	1-cyclopropyl-ethyl		

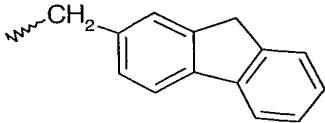
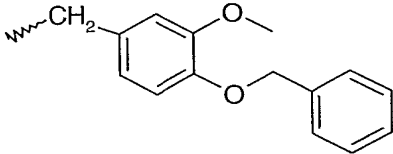
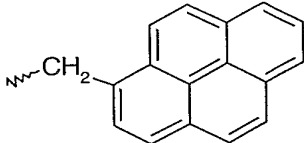
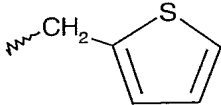
No.	R ₃	Retention time (min.)	
		B1a	B1b
1.072			
1.073			
1.074			
1.075			
1.076			
1.077			
1.078			
1.079	3-methylcyclopentyl		
1.080			
1.081			
1.082			
1.083	(pentafluorophenyl)methyl		
1.084	(2,3,5-trichlorophenyl)methyl		
1.085	(2,3,6-trichlorophenyl)methyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.086	(2,3,4-trifluorophenyl)methyl	2.60	2.48
1.087	(2,6-dichlorophenyl)methyl	2.70	-
1.088	(2,3-dichlorophenyl)methyl	2.70	-
1.089	(2-hydroxy-3,5-dichlorophenyl)methyl		
1.090	(2-chloro-6-fluorophenyl)methyl		
1.091	(2-chloro-6-nitrophenyl)methyl		
1.092	(3-chloro-4-nitrophenyl)methyl		
1.093	(2-chloro-5-nitrophenyl)methyl		
1.094	(2,6-difluorophenyl)methyl	2.50	2.44
1.095	(2,3-difluorophenyl)methyl	2.60	-
1.096	(2-hydroxy-5-bromophenyl)methyl		
1.097	(2-hydroxy-5-chlorophenyl)methyl		
1.098	(3-nitro-4-hydroxyphenyl)methyl		
1.099	(3-hydroxy-4-nitro-phenyl)methyl		
1.100	(3-hydroxy-phenyl)methyl		
1.101	(2-hydroxyphenyl)methyl		
1.102	(2,5-dihydroxyphenyl)methyl		
1.103	3-trifluoromethylcyclohexyl		
1.104			
1.105			
1.106			
1.107	2-methoxycyclohexyl		
1.108			
1.109			

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.110		2.70	2.65
1.111			
1.112			
1.113	(2-trifluoromethylphenyl)methyl		
1.114	(3-trifluoromethoxyphenyl)methyl		
1.115	(3-cyanophenyl)methyl		
1.116	(2,3-methylenedioxyphenyl)methyl		
1.117	(2-methoxy-5-bromo-phenyl)methyl		
1.118	(3-bromo-4-hydroxy-5-methoxy-phenyl)methyl		
1.119	(2-nitro-3-methoxyphenyl)methyl		
1.120	(4-methoxyphenyl)methyl		
1.121	(3-hydroxy-4-methoxyphenyl)methyl		
1.122		2.70	2.65
1.123	2-ethyl-hexyl		
1.124			
1.125	(3,5-bis(trifluoromethyl)-phenyl)methyl		
1.126	(3-pentafluoroethoxyphenyl)methyl		
1.127			
1.128			
1.129	(3-bromo-4,6-dimethoxyphenyl)methyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.130	(2-bromo-4,5-dimethoxyphenyl)methyl		
1.131			
1.132	(2-nitro-4,5-dimethoxyphenyl)methyl		
1.133	2-benzyloxyethyl		
1.134	(3,5-dimethyl-4-hydroxy-phenyl)methyl		
1.135	(2-hydroxy-4,6-dimethoxyphenyl)methyl		
1.136			
1.137			
1.138			
1.139			
1.140	(2,4,6-trimethylphenyl)methyl		
1.141	3,7-dimethyloct-6-en-1-yl		
1.142	naphth-1-yl		
1.143	naphth-2-yl		
1.144			
1.145			

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.146			
1.147			
1.148			
1.149			
1.150			
1.151	n-dodecyl		
1.152	(3-[3,4-dichlorophenoxy]-phenyl)methyl		
1.153	(3-[4-chlorophenoxy]-phenyl)methyl		
1.154			
1.155			

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.156			
1.157			
1.158			
1.159		5.87	5.39
1.160	allyl		
1.161	2-hydroxyethyl		
1.162	2-methyl-n-propyl	2.40	2.28
1.163	n-pentyl	2.57	2.40
1.164	n-hexyl	2.56	2.53
1.165	n-heptyl	2.78	2.61
1.166	n-octyl	2.90	2.69
1.167	n-decyl	3.11	2.94
1.168	2-phenyl-ethyl	2.60	2.40
1.169	trimethylsilyl-methyl	2.69	2.40
1.170	4-acetoxy-n-butyl	2.40	2.19
1.171	3,3,3-trifluoro-n-propyl		
1.172	2-buten-1-yl		
1.173	2-methyl-prop-1-en-3-yl	2.50	2.36
1.174	3-methyl-but-2-en-1-yl		
1.175	2-penten-1-yl		
1.176	3-methoxycarbonyl-(E)-2-propen-1-yl		
1.177	3-ethoxycarbonyl-E-2-propen-1-yl		
1.178	2-methoxycarbonyl-1-propen-3-yl		
1.179	2-ethoxycarbonyl-1-propen-3-yl		
1.180	2-bromo-prop-2-en-1-yl	2.50	2.36

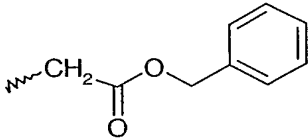
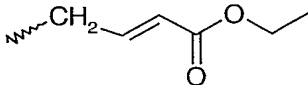
No.	R ₃	Retention time (min.)	
		B1a	B1b
1.181	2-tert-butoxycarbonyl-1-propen-3-yl		
1.182	(E)-3,7-dimethyl-2,6-octadien-1-yl	2.90	2.82
1.183	isobutenyl-oxycarbonyl-methyl	2.57	-
1.184	cyclohexyl-oxycarbonyl-methyl	2.73	-
1.185	benzyl-oxycarbonyl-methyl		
1.186	phenoxyethyl-oxycarbonyl-methyl		
1.187	p-nitrophenyl-oxycarbonyl-methyl		
1.188	(n-butoxycarbonylmethoxy)-carbonyl-methyl	2.69	-
1.189	(o-chlorophenyl)-aminocarbonyl-methyl	2.60	-
1.190	(3,5-dichlorophenyl)-aminocarbonyl-methyl		
1.191	(2-trifluoromethyl-4-nitro-phenyl)-aminocarbonyl-methyl	2.70	-
1.192	(4-methylphenyl)-methyl		
1.193	(3-methylphenyl)-methyl		
1.194	(2-methylphenyl)-methyl		
1.195	(2-fluorophenyl)-methyl		
1.196	(4-cyanophenyl)-methyl	2.50	-
1.197	(2-cyanophenyl)-methyl	2.40	2.36
1.198	(3,5-dimethylphenyl)-methyl		
1.199	(3-methoxyphenyl)-methyl		
1.200	(3-methyl-2-fluorophenyl)-methyl		
1.201	(4-chlorophenyl)-methyl		
1.202	(3-chlorophenyl)-methyl		
1.203	(2-chlorophenyl)-methyl		
1.204	(3,5-difluorophenyl)-methyl		
1.205	(2,4-difluorophenyl)-methyl		
1.206	(3,4-difluorophenyl)-methyl		
1.207	(2,5-difluorophenyl)-methyl		
1.208	(4-nitrophenyl)-methyl		
1.209	(2-nitrophenyl)-methyl		
1.210	2-naphthyl-methyl		
1.211	(3-chloro-2-fluorophenyl)-methyl	2.70	2.53
1.212	(2-chloro-4-fluorophenyl)-methyl		
1.213	(3,4,5-trifluorophenyl)-methyl	2.60	-
1.214	(2,4,6-trifluorophenyl)-methyl		
1.215	(2,4,5-trifluorophenyl)-methyl	2.60	2.48
1.216	(2,3,5-trifluorophenyl)-methyl	2.60	2.44

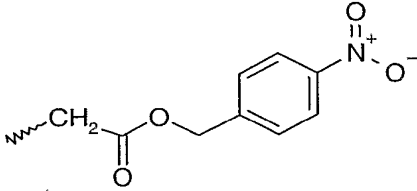
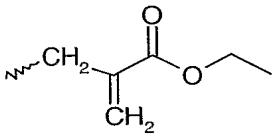
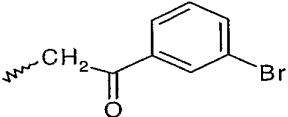
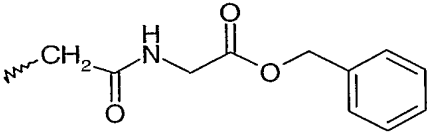
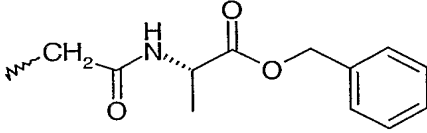
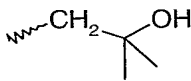
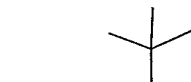
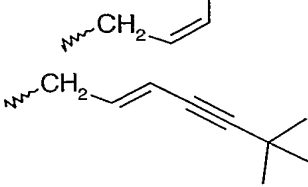
No.	R ₃	Retention time (min.)	
		B1a	B1b
1.217	(4-methoxycarbonylphenyl)-methyl		
1.218	(3-methoxycarbonylphenyl)-methyl		
1.219	(4-difluoromethoxyphenyl)-methyl		
1.220	(3-difluoromethoxyphenyl)-methyl		
1.221	(2-difluoromethoxyphenyl)-methyl		
1.222	(4-trifluoromethylphenyl)-methyl		
1.223	(3-trifluoromethylphenyl)-methyl		
1.224	(3,4-dichlorophenyl)-methyl	2.70	-
1.225	(2,5-dichlorophenyl)-methyl	2.70	-
1.226	(2-(methoxycarbonylmethyl)-phenyl)-methyl		
1.227	(2-methoxy-4-nitrophenyl)-methyl		
1.228	biphenyl-4-yl-methyl	3.60	-
1.229	biphenyl-2-yl-methyl	3.50	-
1.230	(4-methylsulphonylphenyl)-methyl		
1.231	(4-bromophenyl)-methyl		
1.232	(4-trifluoromethoxyphenyl)-methyl		
1.233	(2-fluoro-4-trifluoromethylphenyl)-methyl	2.70	2.57
1.234	(4-fluoro-2-trifluoromethylphenyl)-methyl		
1.235	(4-fluoro-3-trifluoromethylphenyl)-methyl		
1.236	(5-fluoro-2-trifluoromethylphenyl)-methyl	2.70	2.57
1.237	(2-fluoro-5-trifluoromethylphenyl)-methyl	2.70	2.53
1.238	(2-fluoro-3-trifluoromethylphenyl)-methyl		
1.239	(3-fluoro-5-trifluoromethylphenyl)-methyl	2.70	2.57
1.240	(3-fluoro-4-trifluoromethylphenyl)-methyl	2.70	2.57
1.241	(2-methoxy-4-methoxycarbonylphenyl)-methyl		
1.242	(2-fluoro-4-bromophenyl)-methyl		
1.243	(5-fluoro-2-bromophenyl)-methyl	2.70	2.48
1.244	(2,3-dichloro-4-methoxyphenyl)-methyl		
1.245	(4-trifluoromethylthio-phenyl)-methyl		
1.246	(3-trifluoromethylthio-phenyl)-methyl		
1.247	(2-trifluoromethylthio-phenyl)-methyl		
1.248	(4-(o-cyanophenyl)-phenyl)-methyl	3.50	-
1.249	(2-chloro-3-trifluoromethylphenyl)-methyl	2.70	2.61
1.250	(3,4-difluoro-5-trifluorophenyl)-methyl	2.70	-
1.251	(2-nitro-4,5-dimethoxyphenyl)-methyl		
1.252	(4-iodophenyl)-methyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
1.253	(3-iodophenyl)-methyl	2.80	2.65
1.254	(2-iodophenyl)-methyl		
1.255	(2,4-bis(trifluoromethyl)-phenyl)-methyl		
1.256	(3,5-dibromophenyl)-methyl		
1.257	5-bromo-thiophene-2-yl-methyl		
1.258	4-bromo-thiophene-2-yl-methyl		
1.259	10-undecen-1-yl		
1.260	(4-isobutylphenyl)-methyl		
1.261	(2-isopropylthiophenyl)-methyl		
1.262	(4-isopropylphenyl)-methyl		
1.262	(3-isopropylphenyl)-methyl		
1.263	2-ethyl-n-butyl		
1.264	n-nonyl		
1.265	n-undecyl		
1.266	isopentyl		
1.267	(4-ethylphenyl)-methyl		
1.268	(1-methylthio-1-cyclopropyl)-methyl		
1.269	3-methylthio-n-propyl		
1.270	(4-methylthiophenyl)-methyl		
1.271	(2,5-dimethylphenyl)-methyl		
1.272	(3-fluoro-4-methylphenyl)-methyl		
1.273	(5-methyl-thien-2-yl		
1.274	(3-methyl-thien-2-yl		
1.275	(4-methylphenyl)-ethyl		
1.276	(3,5-dichlorophenyl)-methyl		
1.277	(2,4-dichlorophenyl)-methyl		
1.278	1-methyl-1H-pyrrole-2-yl-methyl		
1.279	(3-fluoro-4-chlorophenyl)-methyl		
1.280	(4-fluorophenyl)-methyl		
1.281	4-fluorophenylthio-3-n-propyl		
1.281	cyclopropyl-methyl		
1.282	cyclohexyl-methyl		
1.283	cyclohexyl-3-n-propyl		
1.284	2-pyridyl-methyl		
1.285	3-furyl-methyl		
1.286	3-thienyl-methyl		

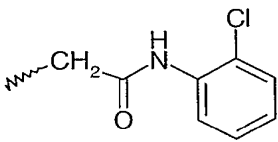
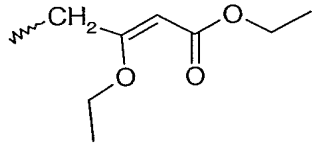
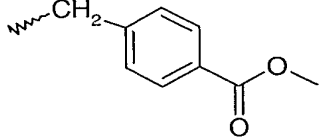
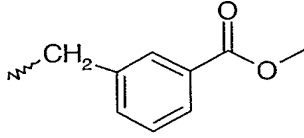
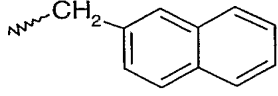
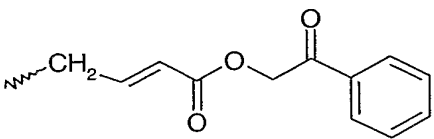
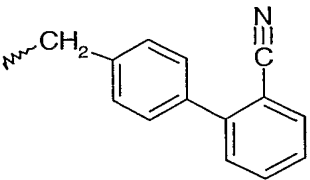
No.	R ₃	Retention time (min.)	
		B1a	B1b
1.287	2-thiazolyl-methyl		
1.288	neopentyl		
1.289	isohexyl		
1.290	tert-butyl		
1.291	3-hydroxypropyl		

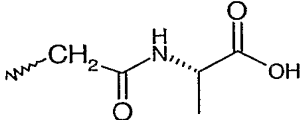
Table 2: Compounds of formula (I) wherein R₁ is sec-butyl (B1a) or isopropyl (B1b) and R₂ is methyl:

No.	R ₃	Retention time (min.)	
		B1a	B1b
2.001	benzyl		
2.002	allyl		
2.003	-CH ₂ -C(=O)-O-ethyl		
2.004	2-hydroxyethyl		
2.005	-CH ₂ -C(=O)-O-methyl		
2.006	but-2-en-1-yl		
2.007			
2.008	-CH ₂ -C(=O)-O-tert-butyl		
2.009	(4-fluoro-2-trifluoromethylphenyl)methyl		
2.010	(4-fluoro-3-trifluoromethylphenyl)methyl		
2.011	(3,4-difluorophenyl)methyl		
2.012	isopropyl		
2.013	(3-trifluoromethylphenyl)methyl		
2.014			
2.015	2-methylallyl		
2.016	-CH ₂ -C(=O)-OH		
2.017	-CH ₂ -C(=O)-NH ₂		

No.	R ₃	Retention time (min.)	
		B1a	B1b
2.018			
2.019	(2-methylphenyl)methyl		
2.020			
2.021			
2.022	ethyl		
2.023			
2.024			
2.025			
2.026	(4-phenylphenyl)methyl		
2.027			
2.028			
2.029	(4-tert-butylphenyl)methyl		
2.030	(4-fluorophenyl)methyl		
2.031	(4-bromophenyl)methyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
2.032	2-bromoallyl		
2.033	-CH ₂ -C(=O)-O-isopropyl		
2.034	-CH ₂ -C(=O)-O-CH ₂ -CH ₂ -O-CH ₃		
2.035	(2,3,4-trifluorophenyl)methyl		
2.036	(2,4,5-trifluorophenyl)methyl		
2.037	(2,3,6-trifluorophenyl)methyl		
2.038	(3,5-dibromophenyl)methyl		
2.039	(3-fluoro-6-bromophenyl)methyl		
2.040	(2,3-dichlorophenyl)methyl		
2.041	(2,6-dichlorophenyl)methyl		
2.042	(2,5-dichlorophenyl)methyl		
2.043	(3,4-dichlorophenyl)methyl		
2.044	(2-fluoro-3-chloro-phenyl)methyl		
2.045	(2-chloro-4-fluoro-phenyl)methyl		
2.046	(2,5-difluorophenyl)methyl		
2.047	(2,6-difluorophenyl)methyl		
2.048	(2,3-difluorophenyl)methyl		
2.049	(3,5-difluorophenyl)methyl		
2.050	(2-bromophenyl)methyl		
2.051	(2-chlorophenyl)methyl		
2.052	(2-fluorophenyl)methyl		
2.053	(3-fluorophenyl)methyl		
2.054	(4-iodophenyl)methyl		
2.055	(2-iodophenyl)methyl		
2.056	(4-nitrophenyl)methyl		
2.057	(3-nitrophenyl)methyl		
2.058	(3-trifluoromethyl-6-chlorophenyl)methyl		
2.059	(3-trifluoromethyl-5-fluorophenyl)methyl		
2.060	(3,5-dichlorophenyl)methyl		
2.061	(2-trifluoromethylphenyl)methyl		
2.062	(3-trifluoromethoxyphenyl)methyl		
2.063	(4-cyanophenyl)methyl		
2.064	(2-cyanophenyl)methyl		
2.065	(3-cyanophenyl)methyl		
2.066	(2,3-dichloro-4-methoxy-phenyl)methyl		

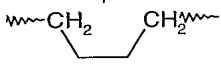
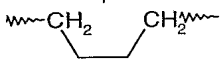
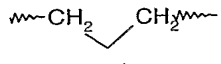
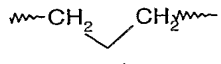
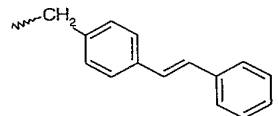
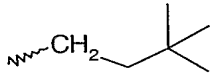
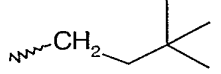
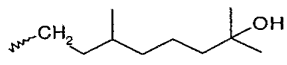
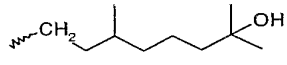
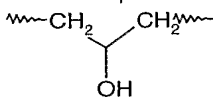
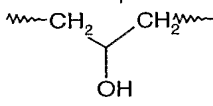
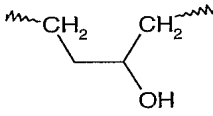
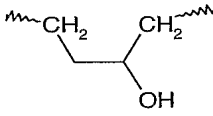
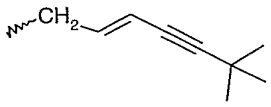
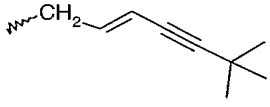
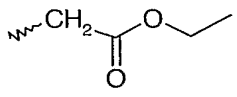
No.	R ₃	Retention time (min.)	
		B1a	B1b
2.067			
2.068	(4-difluoromethoxyphenyl)methyl		
2.069	(2-fluoro-3-methylphenyl)methyl		
2.070			
2.071	(3,5-bis(trifluoromethyl)-phenyl)methyl		
2.072	(2,4-bis(trifluoromethyl)-phenyl)methyl		
2.073			
2.074			
2.075	(3,5-dimethylphenyl)methyl		
2.076	(2-methoxy-4-methoxycarbonyl-phenyl)methyl		
2.077			
2.078			
2.079	(2-phenylphenyl)methyl		
2.080	(4-phenylphenyl)methyl		
2.081			
2.082	n-propyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
2.083	isopropyl		
2.084	n-butyl		
2.085	n-pentyl		
2.086	n-hexyl		
2.087	n-heptyl		
2.088	n-octyl		
2.089	isobutyl		
2.090	sec.-butyl		
2.091	tert.-butyl		
2.092	isopentyl		
2.093	neopentyl		
2.094	isohexyl		
2.095			
2.096	(4-methylphenyl)-methyl		
2.097	(3-methylphenyl)-methyl		
2.098	(3-methoxyphenyl)-methyl		
2.099	(4-chlorophenyl)-methyl		
2.100	(3-chlorophenyl)-methyl		
2.101	(2,4-difluorophenyl)-methyl		
2.102	(2-nitrophenyl)-methyl		
2.103	(3,4,5-trifluorophenyl)-methyl		
2.104	(2,4,6-trifluorophenyl)-methyl		
2.105	(2,3,5-trifluorophenyl)-methyl		
2.106	2-penten-1-yl		
2.107	3,3-dimethyl-allyl		
2.108	(3-difluoromethoxyphenyl)-methyl		
2.109	(2-difluoromethoxyphenyl)-methyl		
2.110	(4-trifluoromethylphenyl)-methyl		
2.111	(3-methoxycarbonylmethylphenyl)-methyl		
2.112	(4-methylsulphonylphenyl)-methyl		
2.113	(3-bromophenyl)-methyl		
2.114	(4-trifluoromethoxyphenyl)-methyl		
2.115	(2-fluoro-4-trifluoromethylphenyl)-methyl		

No.	R ₃	Retention time (min.)	
		B1a	B1b
2.116	(5-fluoro-2-trifluoromethylphenyl)-methyl		
2.117	(2-fluoro-5-trifluoromethylphenyl)-methyl		
2.118	(2-fluoro-3-trifluoromethylphenyl)-methyl		
2.119	(3-fluoro-4-trifluoromethylphenyl)-methyl		
2.120	(2,3,6-trichlorophenyl)-methyl		
2.121	pentafluorophenyl-methyl		
2.122	(4-bromo-2-fluorophenyl)-methyl		
2.123	(3-trifluoromethylthiophenyl)-methyl		
2.124	(4-trifluoromethylthiophenyl)-methyl		
2.125	(2-trifluoromethylthiophenyl)-methyl		
2.126	(3,4-difluoro-5-trifluoromethylphenyl)-methyl		
2.127	(3-iodophenyl)-methyl		
2.128	(5-bromo-thien-2-yl)-methyl		
2.129	(4-bromo-thien-2-yl)-methyl		
2.130	10-undecen-1-yl		
2.131	(4-isobutylphenyl)-methyl		
2.132	(2-isopropylthiophenyl)-methyl		
2.133	(4-isopropylphenyl)-methyl		
2.134	(3-isopropylphenyl)-methyl		
2.135	2-ethyl-n-butyl		
2.136	4-decen-1-yl		
2.137	n-nonyl		
2.138	n-decyl		
2.139	n-undecyl		
2.140	n-dodecyl		
2.141	(4-ethylphenyl)-methyl		
2.142	3-methylthio-n-propyl		
2.143	([4-methylthio]phenyl)-methyl		
2.144	(2,4,6-trimethylphenyl)-methyl		
2.145	(2,5-dimethylphenyl)-methyl		
2.146	(2,4-dimethylphenyl)-methyl		
2.147	(3-fluoro-4-methylphenyl)-methyl		
2.148	(5-methyl-furan-2-yl)-methyl		
2.149	(5-methyl-thien-2-yl)-methyl		
2.150	(4-methylphenyl)-3-ethyl		
2.151	(3-methyl-thien-2-yl)-methyl		

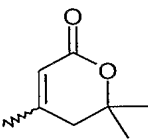
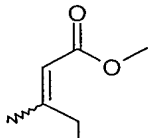
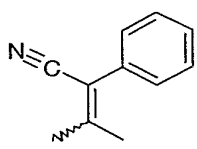
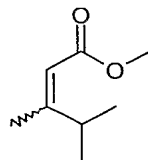
No.	R ₃	Retention time (min.)	
		B1a	B1b
2.152	(2,3,5-trichlorophenyl)-methyl		
2.153	(2,3,6-trichlorophenyl)-methyl		
2.154	(2,4-dichlorophenyl)-methyl		
2.155	(3-fluoro-4-chlorophenyl)-methyl		
2.156	(3,4-difluorophenyl)-methyl		
2.157	Cyclopropylmethyl		
2.158	cyclohex-3-en-1-yl-methyl		
2.159	cyclohexyl-methyl		
2.160	cyclohexyl-n-propyl		
2.161	phenyl-3-n-propyl		
2.162	phenyl-2-ethyl		
2.163	2-pyridyl-methyl		
2.164	3-pyridyl-methyl		
2.165	2-furyl-methyl		
2.166	3-furyl-methyl		
2.167	4-pyridyl-methyl		
2.168	3-thienyl-methyl		
2.169	2-thiazolyl-methyl		
2.170	isobutenyl-oxycarbonyl-methyl		
2.171	cyclohexyl-oxycarbonyl-methyl		
2.172	n-butoxycarbonyl-methoxycarbonyl-methyl		
2.173	phenoxyethyl-oxycarbonyl-methyl		
2.174	p-nitrophenyl-oxycarbonyl-methyl		
2.175	3-phenyl-allyl		
2.176	3,5-dichlorophenyl-aminocarbonyl-methyl		
2.177	2-(tert-butoxycarbonyl)-allyl		
2.178	4-acetoxy-n-butyl		
2.179	2-trifluoromethyl-ethyl		
2.180	trimethylsilyl-methyl		
2.181	3,7-dimethyloct-6-en-1-yl		
2.182	3-hydroxypropyl		

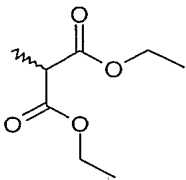
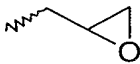
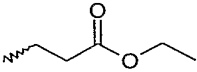
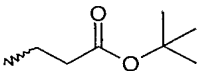
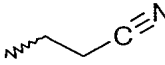
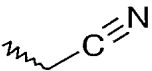
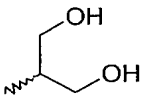
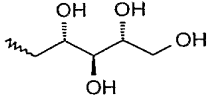
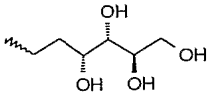
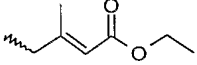
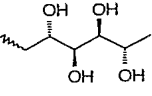
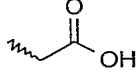
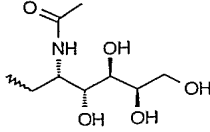
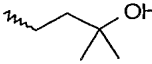
Table 3: Compounds of formula (I) wherein R₁ is sec-butyl (B1a) or isopropyl (B1b):

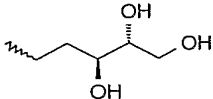
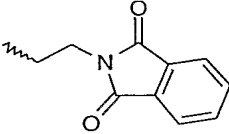
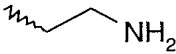
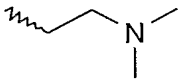
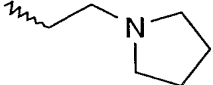
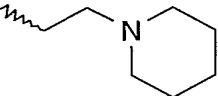
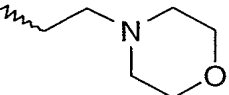
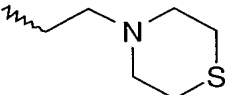
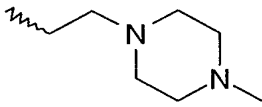
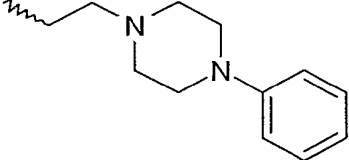
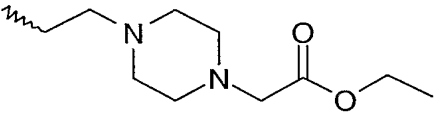
No.	R ₂	R ₃	Retention time (min.)	
			B1a	B1b
3.001	3-phenylallyl	3-phenylallyl		
3.002				
3.003				
3.004	3-phenyl-n-propyl	3-phenyl-n-propyl		
3.005	3,7-dimethyloct-6-en-1-yl	3,7-dimethyloct-6-en-1-yl		
3.006	ethyl			
3.007	ethyl	ethyl		
3.008				
3.009				
3.010				
3.011				
3.012				
3.013	ethyl	benzyl		
3.014	ethyl	allyl		
3.015	ethyl			
3.016	3-methyl-n-butyl	3-methyl-n-butyl		
3.017	3-phenyl-n-butyl	3-phenyl-n-butyl		
3.018	ethyl	n-propyl		
3.019	ethyl	isopropyl		
3.020	ethyl	n-butyl		
3.021	ethyl	pentyl		

No.	R ₂	R ₃	Retention time (min.)	
			B1a	B1b
3.022	ethyl	hexyl		
3.023	ethyl	heptyl		
3.024	ethyl	n-octyl		
3.025	ethyl	sec-butyl		
3.026	ethyl	tert-butyl		
3.027	ethyl	isopentyl		
3.028	ethyl	neopentyl		
3.029	ethyl	isohexyl		

Table 4: Compounds of formula (I) wherein R₁ is sec-butyl (B1a) or isopropyl (B1b) and R₂ is hydrogen:

No.	R ₃	Retention time (min.)	
		B1a	B1b
4.001			
4.002			
4.003			
4.004			

No.	R ₃	Retention time (min.)	
		B1a	B1b
4.005			
4.006			
4.007			
4.008			
4.009			
4.010			
4.011			
4.012			
4.013			
4.014			
4.015			
4.016			
4.017			
4.018			

No.	R ₃	Retention time (min.)	
		B1a	B1b
4.019			
4.020			
4.021			
4.022			
4.023			
4.024			
4.025			
4.026			
4.027			
4.028			
4.029			

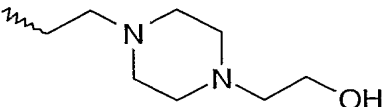
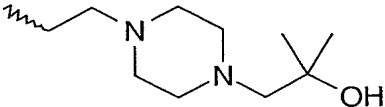
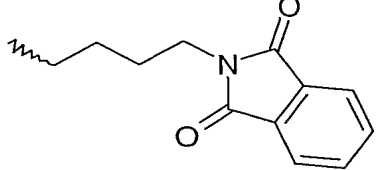
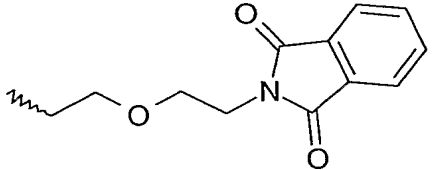
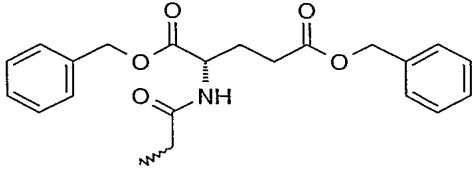
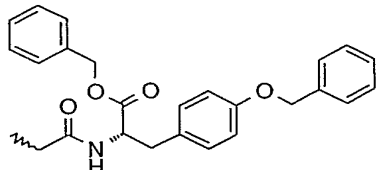
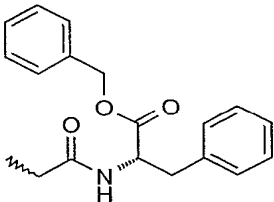
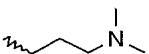
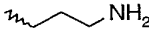
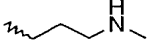
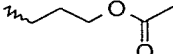

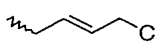
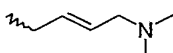
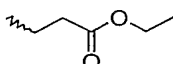
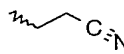
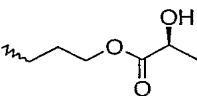
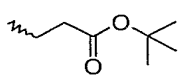


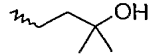
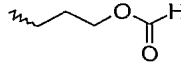
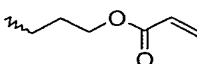
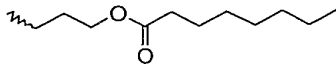
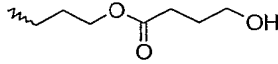
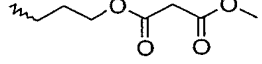
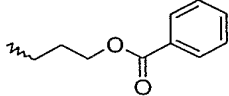
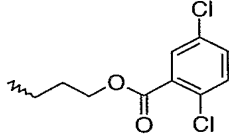
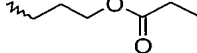
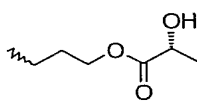
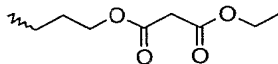
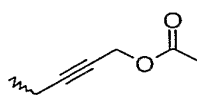
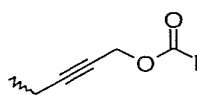
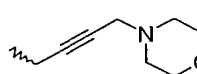
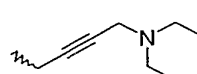

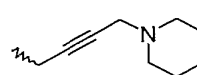
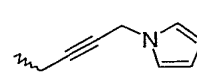
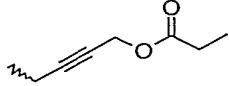
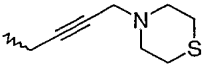
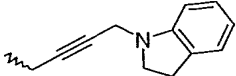
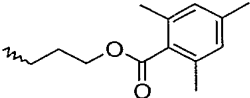
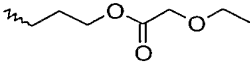
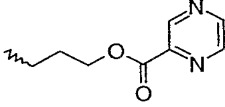
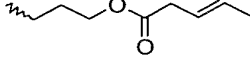
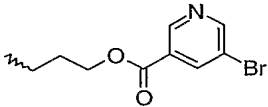
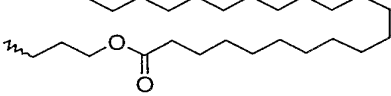
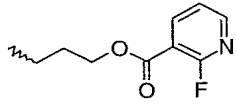
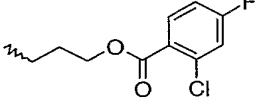
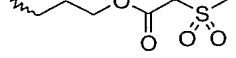
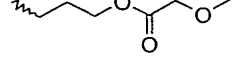
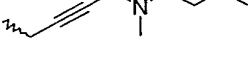
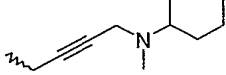
No.	R ₃	Retention time (min.)	
		B1a	B1b
4.030			
4.031			
4.032			
4.033			

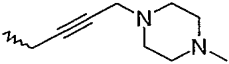
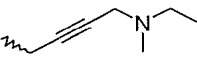
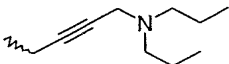
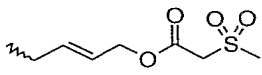
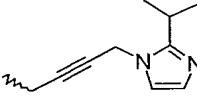
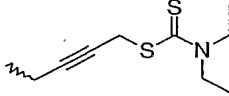
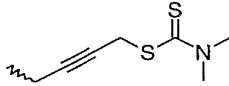
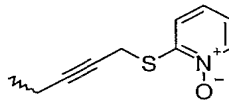
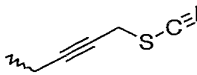
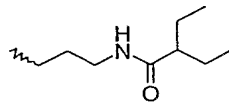
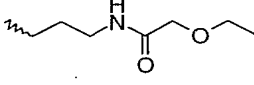
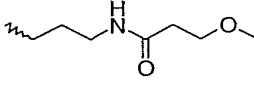
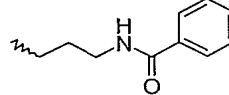
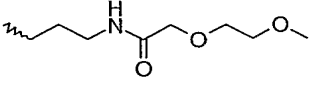
Table 5: Compounds of formula (I) wherein R₁ is sec-butyl (B1a) or isopropyl (B1b) and R₂ is methyl

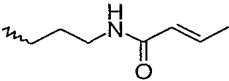
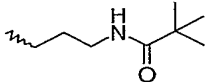
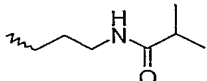
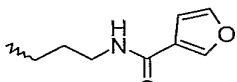
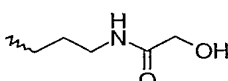
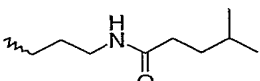
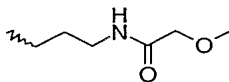
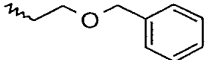
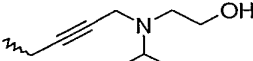
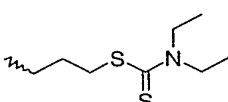
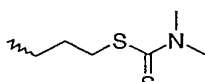
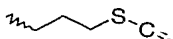
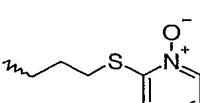


No.	R ₃	Retention time (min.)	
		B1a	B1b
5.001 5.002	trifluoromethyl 		
5.003 5.004 5.005	3,3-dichloroallyl 2-chloroallyl 		

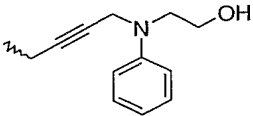
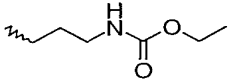
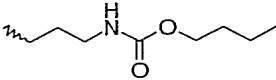
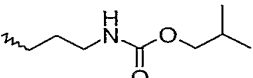
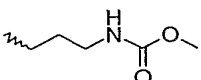
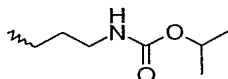
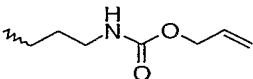
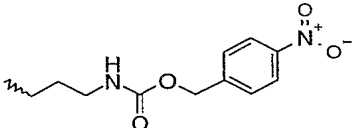
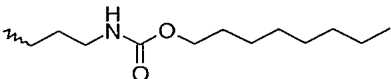
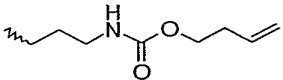
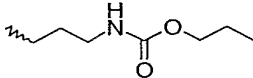
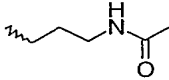
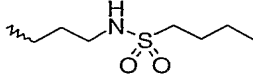
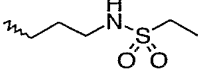
No.	R ₃	Retention time (min.)	
		B1a	B1b
5.006			
5.007	3-iodopropyl		
5.008	cyclobutyl		
5.009			
5.010			
5.011			
5.012			
5.013			
5.014			
5.015			
5.016			
5.017			
5.018			
5.019			
5.020			
5.021			
5.022			
5.023			
5.024			

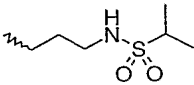
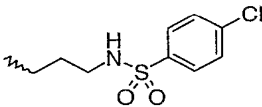
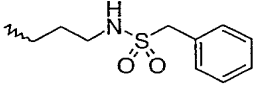
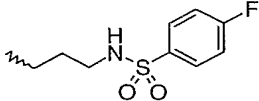
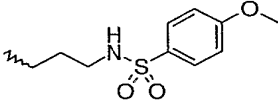
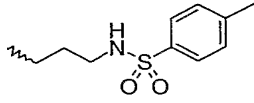
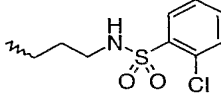
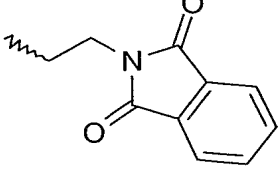
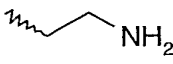
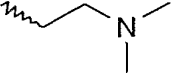
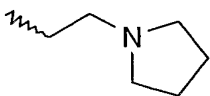
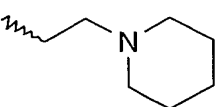
No.	R ₃	Retention time (min.)	
		B1a	B1b
5.025			
5.026			
5.027			
5.028			
5.029			
5.030			
5.031			
5.032			
5.033			
5.034			
5.035			
5.036			
5.037			
5.038			
5.039			
5.040			

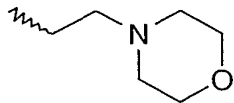
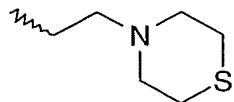
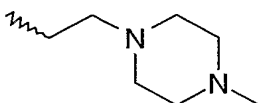
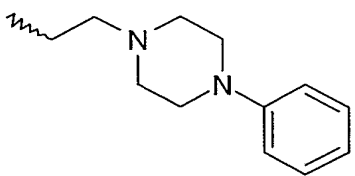
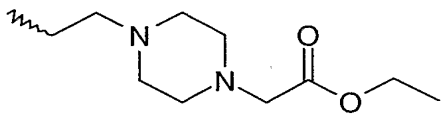
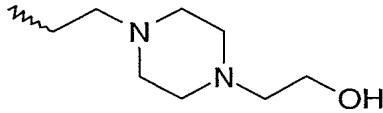
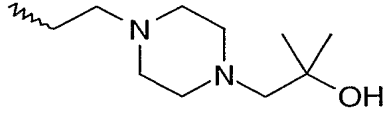
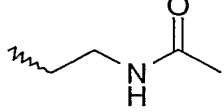
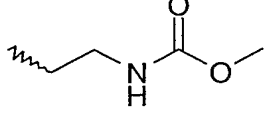
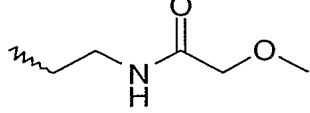
No.	R ₃	Retention time (min.)	
		B1a	B1b
5.041			
5.042			
5.043			
5.044			
5.045			
5.046			
5.047			
5.048			
5.049			
5.050			
5.051			
5.052			
5.053			
5.054			

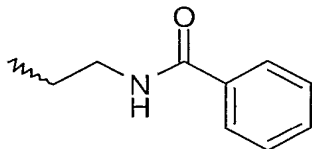
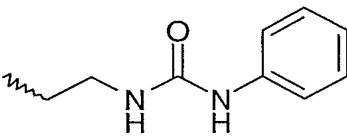
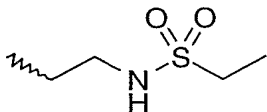
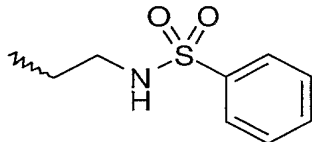
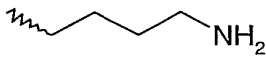
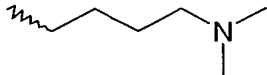
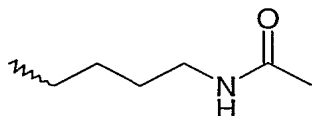
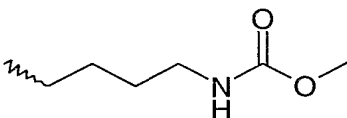
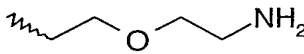
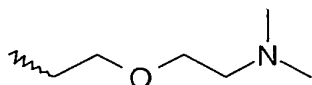
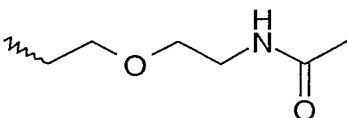
No.	R ₃	Retention time (min.)	
		B1a	B1b
5.055			
5.056			
5.057			
5.058			
5.059			
5.060			
5.061			
5.062			
5.063			
5.064			
5.065			
5.066			
5.067			
5.068			

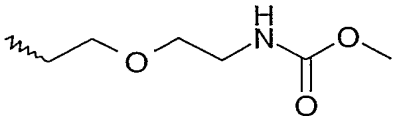
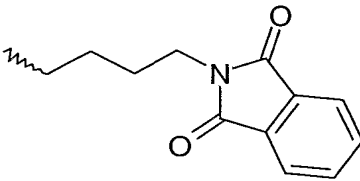
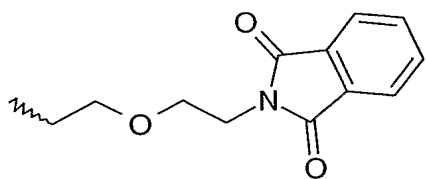
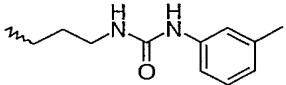
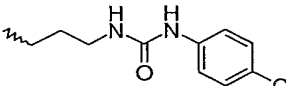
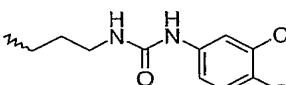
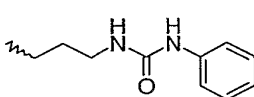
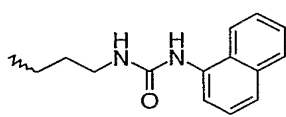
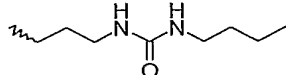
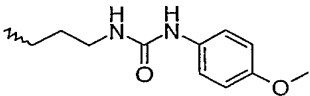
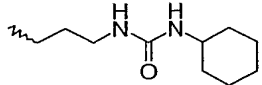
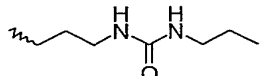
No.	R ₃	Retention time (min.)	
		B1a	B1b
5.069			
5.070			
5.071			
5.072			
5.073			
5.074			
5.075			
5.076			
5.077			
5.078			
5.079			
5.080			
5.081			
5.082			
5.083			

No.	R ₃	Retention time (min.)	
		B1a	B1b
5.084			
5.085			
5.086			
5.087			
5.088			
5.089			
5.090			
5.091			
5.092			
5.093			
5.094			
5.095			
5.096			
5.097			

No.	R ₃	Retention time (min.)	
		B1a	B1b
5.098			
5.099			
5.100			
5.101			
5.102			
5.103			
5.104			
5.105			
5.106			
5.108			
5.109			
5.110			

No.	R ₃	Retention time (min.)	
		B1a	B1b
5.111			
5.112			
5.113			
5.114			
5.115			
5.116			
5.117			
5.118			
5.119			
5.120			

No.	R ₃	Retention time (min.)	
		B1a	B1b
5.121			
5.122			
5.123			
5.124			
5.125			
5.126			
5.127			
5.128			
5.129			
5.130			
5.131			

No.	R ₃	Retention time (min.)	
		B1a	B1b
5.132			
5.133			
5.134			
5.135			
5.136			
5.137			
5.138			
5.139			
5.140			
5.141			
5.142			
5.143			

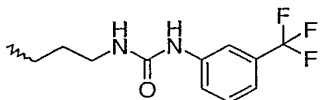
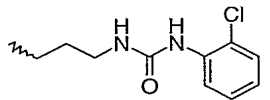
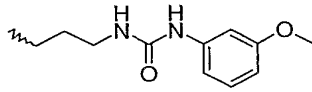
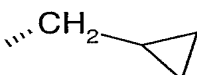
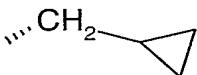
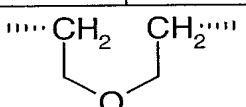
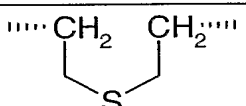
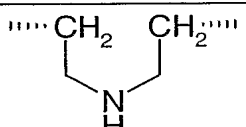
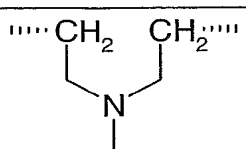
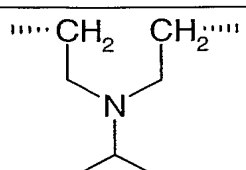
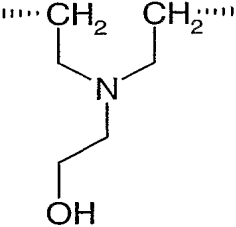
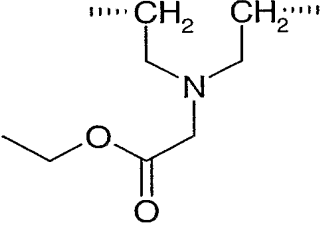
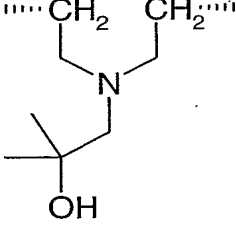
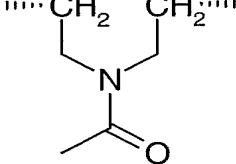
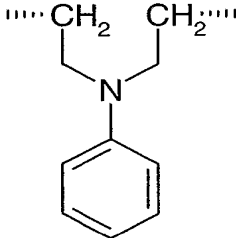
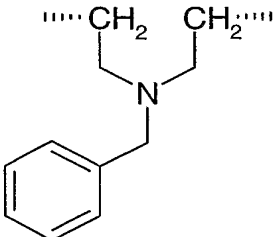
No.	R ₃	Retention time (min.)	
		B1a	B1b
5.144			
5.145			
5.146			

Table 6: Compounds of formula (I) wherein R₁ is sec-butyl (B1a) or isopropyl (B1b)

No.	R ₂	R ₃	Retention time (min.)	
			B1a	B1b
6.001			6.13	5.71
6.002				
6.003				
6.004				
6.005				
6.007				

No.	R ₂	R ₃	Retention time (min.)	
			B1a	B1b
6.008				
6.009				
6.010				
6.011				
6.012				
6.013				

Formulation Examples for use in crop protection (% = percent by weight)

<u>Example F1: Emulsifiable concentrates</u>	a)	b)	c)
active ingredient	25 %	40 %	50 %
calcium dodecylbenzenesulfonate	5 %	8 %	6 %
castor oil polyethylene glycol ether (36 mol of ethylene oxide)	5 %	-	-
tributylphenol polyethylene glycol ether (30 mol of ethylene oxide)	-	12 %	4 %
cyclohexanone	-	15 %	20 %
xylene mixture	65 %	25 %	20 %

Mixing together the finely ground active ingredient and additives yields an emulsifiable concentrate which, on dilution with water, yields emulsions of the desired concentration.

<u>Example F2: Solutions</u>	a)	b)	c)	d)
active ingredient	80 %	10 %	5 %	95 %
ethylene glycol monomethyl ether		20 %	-	-
polyethylene glycol (MW 400)		70 %	-	-
N-methyl-2-pyrrolidone	20 %	20 %	1%	5%
epoxidised coconut oil	-	-	-	1 %
petroleum ethers (boiling range 160-190°)	-	-	94 %	-

Mixing together the finely ground active ingredient and additives yields a solution suitable for application in the form of microdrops.

<u>Example F3: Granules</u>	a)	b)	c)	d)
active ingredient	5 %	10 %	8 %	21 %
kaolin	94 %	-	79 %	54 %
highly disperse silicic acid	1 %	-	13 %	7 %
attapulgite	-	90 %	-	18 %

The active ingredient is dissolved in dichloromethane, the solution is sprayed onto the carrier mixture, and the solvent is evaporated off *in vacuo*.

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Example F4: Wettable powders

	a)	b)	c)
active ingredient	25 %	50 %	75 %
sodium lignosulfonate	5 %	5 %	-
sodium lauryl sulfate	3 %	-	5 %
sodium diisobutyl-naphthalene-sulfonate	-	6 %	10 %
octylphenol polyethylene glycol ether (7-8 mol of ethylene oxide)	-	2 %	-
highly disperse silicic acid	5 %	10 %	10 %
kaolin	62 %	27 %	-

Active ingredient and additives are mixed together and the mixture is ground in a suitable mill, yielding wettable powders which can be diluted with water to give suspensions of the desired concentration.

Example F5: Emulsifiable concentrate

active ingredient	10 %
octylphenol polyethylene glycol ether (4-5 mol of ethylene oxide)	3 %
calcium dodecylbenzenesulfonate	3 %
castor oil polyethylene glycol ether (36 mol of ethylene oxide)	4 %
cyclohexanone	30 %
xylene mixture	50 %

Mixing together the finely ground active ingredient and additives yields an emulsifiable concentrate which, on dilution with water, yields emulsions of the desired concentration.

Example F6: Extruder granules

active ingredient	10 %
sodium lignosulfonate	2 %
carboxymethylcellulose	1 %
kaolin	87 %

Active ingredient and additives are mixed together and the mixture is ground, moistened with water, extruded and granulated, and the granules are dried in a stream of air.

Example F7: Coated granules

active ingredient	3 %
polyethylene glycol (MW 200)	3 %
kaolin	94 %

Uniform application of the finely ground active ingredient to the kaolin moistened with polyethylene glycol in a mixer yields non-dusty coated granules.

Example F8: Suspension concentrate

active ingredient	40 %
ethylene glycol	10 %
nonylphenol polyethylene glycol ether (15 mol of ethylene oxide)	6 %
sodium lignosulfonate	10 %
carboxymethylcellulose	1 %
aqueous formaldehyde solution (37 %)	0.2 %
aqueous silicone oil emulsion (75 %)	0.8 %
water	32 %

Mixing together the finely ground active ingredient and additives yields a suspension concentrate which, on dilution with water, yields suspensions of the desired concentration.

Biological Examples:Example B1: Action against *Spodoptera littoralis*

Young soybean plants are sprayed with an aqueous emulsion spray mixture comprising 12.5 ppm of active ingredient and, after the spray-coating has dried, are populated with 10 caterpillars of *Spodoptera littoralis* in the first stage and then placed in a plastics container. 3 days later, the percentage reduction in population and the percentage reduction in feeding damage (% activity) are determined by comparing the number of dead caterpillars and the feeding damage on the treated plants with that on untreated plants.

The compounds of the Tables exhibit good activity in this test. For example, especially compounds A.2, A.4, A.9, A.10, A.12, A.13, A.14, A.19, A.21 to A.25 and A.28 have activity of more than 90 % in this test.

Example B2: Systemic action against *Spodoptera littoralis*:

Maize seedlings are laid in the solution under test. 6 days later, the leaves are cut off and placed in a petri dish on moist filter paper and infested with from 12 to 15 larvae of

Spodoptera littoralis in the L₁ stage. 4 days later, the percentage reduction in population (% activity) is determined by comparing the number of dead caterpillars on the treated plants with that on untreated plants.

The compounds of the Tables exhibit good activity in this test. For example, especially compounds A.5, A.9, A.10, A.13, A.14, A.19, A.22 to A.25 and A.28 have activity of more than 90 % in this test.

Example B3: Action against *Heliothis virescens*

30-35 eggs of *Heliothis virescens* that are from 0 to 24 hours old are deposited on filter paper lying in a petri dish on a layer of artificial nutrient. 0.8 ml of the solution under test is then pipetted onto the filter papers. Evaluation is carried out after 6 days. The percentage reduction in population (% activity) is determined by comparing the number of dead eggs and larvae on the filter papers with that on untreated filter papers.

The compounds of the Tables exhibit good activity in this test. For example, especially compounds A.2 and A.14 have activity of more than 90 % in this test.

Example B4: Action against *Plutella xylostella* caterpillars

Young cabbage plants are sprayed with an aqueous emulsion spray mixture comprising 12.5 ppm of active ingredient and, after the spray-coating has dried, are populated with 10 caterpillars of *Plutella xylostella* in the first stage and then placed in a plastics container. Evaluation is carried out 3 days later. The percentage reduction in population and the percentage reduction in feeding damage (% activity) are determined by comparing the number of dead caterpillars and the feeding damage on the treated plants with that on untreated plants.

The compounds of Table 1 exhibit good activity against *Plutella xylostella* in this test. For example, especially compounds A.2, A.4, A.5, A.9, A.10, A.12, A.13, A.14, A.19, A.21 to A.25 and A.28 have activity of more than 90 % in this test.

Example B5: Action against *Diabrotica balteata*

Maize seedlings are sprayed with an aqueous emulsion spray mixture comprising 12.5 ppm of active ingredient and, after the spray-coating has dried, are populated with 10 *Diabrotica balteata* larvae in the second stage and then placed in a plastics container. 6 days later, the percentage reduction in population (% activity) is determined by comparing the number of dead larvae on the treated plants with that on untreated plants.

The compounds of the Tables exhibit good activity in this test. For example, especially compounds A.2, A.4, A.9, A.10, A.13, A.14, A.19 and A.23 have activity of more than 90 % in this test.

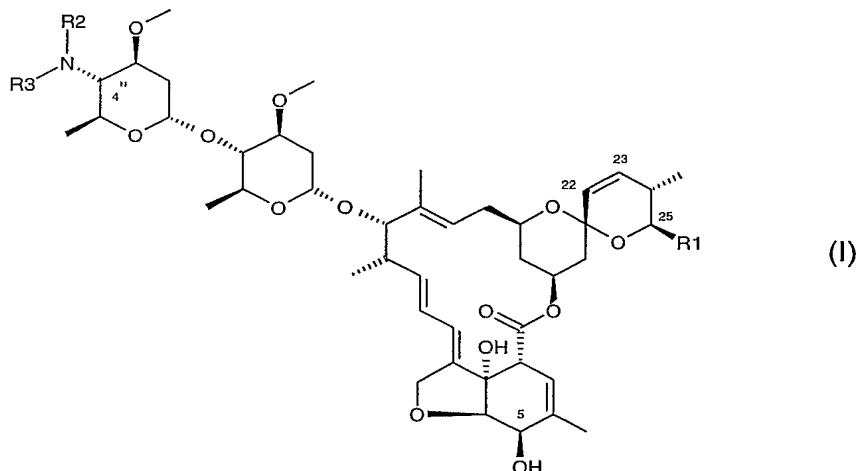
Example B6: Action against *Tetranychus urticae*

Young bean plants are populated with a mixed population of *Tetranychus urticae* and sprayed 1 day later with an aqueous emulsion spray mixture comprising 12.5 ppm of active ingredient, incubated for 6 days at 25°C and then evaluated. The percentage reduction in population (% activity) is determined by comparing the number of dead eggs, larvae and adults on the treated plants with that on untreated plants.

The compounds of the Tables exhibit good activity in this test. For example, especially compounds A.2, A.4, A.5, A.9, A.10, A.13, A.14, A.19, A.21 to A.25 and A.28 have activity of more than 90 % in this test.

WHAT IS CLAIMED IS:

1. A compound of formula



which in the 4"-position has the (*S*)-configuration and wherein

R_1 is C_1 - C_{12} alkyl, C_3 - C_8 cycloalkyl; or C_2 - C_{12} alkenyl;

R_2 is hydrogen, unsubstituted or mono- to penta-substituted C_1 - C_{12} alkyl or unsubstituted or mono- to penta-substituted C_2 - C_{12} alkenyl;

R_3 is C_1 - C_{12} alkyl, mono- to penta-substituted C_1 - C_{12} alkyl, unsubstituted or mono- to penta-substituted C_1 - C_{12} alkoxy- C_1 - C_{12} alkyl, unsubstituted or mono- to penta-substituted C_3 - C_{12} cycloalkyl, unsubstituted or mono- to penta-substituted C_2 - C_{12} alkenyl; unsubstituted or mono- to penta-substituted C_4 - C_{12} cycloalkenyl, unsubstituted or mono- to penta-substituted C_2 - C_{12} alkynyl; or

R_2 and R_3 together are a three- to seven-membered alkylene or four- to seven-membered alkenylene bridge in each of which a CH_2 group may have been replaced by O, S or NR_4 ;

wherein the substituents of the mentioned alkyl, alkoxy-alkyl, alkenyl, alkynyl, alkylene, alkenylene, cycloalkyl and cycloalkenyl radicals are selected from the group consisting of OH, halogen, halo- C_1 - C_2 alkyl, CN, SCN, NO_2 , $Si(C_1-C_{12}alkyl)_3$, C_2 - C_6 alkynyl, C_3 - C_8 cycloalkyl that is unsubstituted or substituted by from one to three methyl groups, norbornylenyl, C_3 - C_8 cycloalkenyl that is unsubstituted or substituted by from one to three methyl groups; C_3 - C_8 halocycloalkyl, C_1 - C_{12} alkoxy, C_1 - C_6 alkoxy- C_1 - C_6 alkoxy, C_3 - C_8 cycloalkoxy, C_1 - C_{12} haloalkoxy, C_1 - C_{12} alkylthio, C_3 - C_8 cycloalkylthio, C_1 - C_{12} haloalkylthio, C_1 - C_{12} alkylsulfinyl,

C₃-C₈cycloalkylsulfinyl, C₁-C₁₂haloalkylsulfinyl, C₃-C₈halocycloalkylsulfinyl, C₁-C₁₂alkylsulfonyl, C₃-C₈cycloalkylsulfonyl, C₁-C₁₂haloalkylsulfonyl, C₃-C₈halocycloalkylsulfonyl, C₂-C₈alkenyl, C₂-C₈alkynyl, NH₂, NH(C₁-C₆alkyl), NH(hydroxy-C₁-C₆alkyl), N(C₁-C₆alkyl)(hydroxy-C₁-C₆alkyl), N(phenyl)(hydroxy-C₁-C₆alkyl), N(C₁-C₆alkyl)₂, -C(=X)ZR₅, -O-C(=X)R₆, -O-C(=X)N(R₈)R₆, -NHC(=X)R₆, -NHC(=X)OR₆, -NHC(=X)SR₆, -NHC(=X)N(R₈)R₆, -S-C(=S)R₆, -NHS(O)₂-R₉, -P(=O)(OC₁-C₆alkyl)₂;

aryl, heterocyclyl, aryloxy, arylthio, heterocyclyloxy; and aryl, heterocyclyl, aryloxy, arylthio and heterocyclyloxy which, depending on the possibilities for substitution on the ring, are mono- to penta-substituted by substituents selected from the group consisting of OH, =O, halogen, CN, NO₂, C₁-C₁₂alkyl, hydroxyC₁-C₁₂alkyl, C₃-C₈cycloalkyl, C₁-C₁₂haloalkyl, C₁-C₁₂alkoxy, C₁-C₁₂haloalkoxy, C₁-C₁₂alkylthio, C₁-C₁₂haloalkylthio, C₁-C₆alkoxy-C₁-C₆alkyl, dimethylamino-C₁-C₆alkoxy, C₂-C₈alkenyl, C₂-C₈alkynyl, Si(C₁-C₁₂alkyl)₃, methylenedioxy, -C(=X)R₅, -O-C(=X)R₆, -CH₂-C(=O)R₅, -CH₂-O-C(=O)R₆, -NH-C(=X)R₆, -S-C(=S)R₆, NH₂, NH(C₁-C₁₂alkyl), -N(C₁-C₁₂alkyl)₂, C₁-C₆alkylsulfinyl, C₃-C₈cycloalkylsulfinyl, C₁-C₆haloalkylsulfinyl, C₃-C₈halocycloalkylsulfinyl, C₁-C₆alkylsulfonyl, C₃-C₈cycloalkylsulfonyl, C₁-C₆haloalkylsulfonyl and C₃-C₈halocycloalkylsulfonyl; unsubstituted phenyl, phenoxy, phenyl-C₁-C₆alkyl, phenyl-C₁-C₆alkoxy, phenyl-C₂-C₆alkenyl, phenyl-C₂-C₆alkynyl; phenyl, phenoxy, phenyl-C₁-C₆alkyl, phenyl-C₁-C₆alkoxy, phenyl-C₂-C₆alkenyl and phenyl-C₂-C₆alkynyl, each of which is substituted in the phenylring by from one to three substituents independently selected of one another from nitro, cyano, halogen, C₁-C₁₂alkoxy, haloC₁-C₁₂alkyl and haloC₁-C₁₂alkoxy;

X is O or S;

Z is a bond, O, NR₈ or S;

R₄ is C₁-C₈alkyl, C₃-C₈cycloalkyl, C₂-C₈alkenyl, C₂-C₈alkynyl, benzyl or -C(=O)-R₅;

R₅ is H, OH, SH, C₁-C₁₂alkyl, C₂-C₈alkenyl, C₂-C₈alkynyl, C₁-C₁₂haloalkyl, C₂-C₈haloalkenyl, C₁-C₆alkoxy-C₁-C₆alkyl, -C₁-C₆S(O)₂-R₉, C₁-C₆alkyl which is mono- or disubstituted with substituents selected from COOH, -C(=O)O-CH₂-phenyl, unsubstituted or substituted phenyl, unsubstituted or substituted biphenyl, wherein the substituents are selected from cyano, halogen, nitro, trifluoromethyl, trifluoromethyl and benzyloxy; aryl, heterocyclyl, aryl-C₁-C₁₂alkyl, aryloxy-C₁-C₁₂alkyl, -C₁-C₆alkyl-C(=O)-R₇, -C₁-C₆alkyl-C(=O)-R₇; or aryl, heterocyclyl, aryl-C₁-C₁₂alkyl, aryloxy-C₁-C₁₂alkyl, each of which is substituted on the aromatic ring by from one to three substituents selected

independently of one another from halogen, nitro, cyano, C₁-C₆alkoxy, C₁-C₆haloalkyl and C₁-C₆haloalkoxy;

R₆ is H, C₁-C₂₄alkyl, C₁-C₆alkoxy-C₁-C₆alkyl, C₁-C₁₂hydroxyalkyl, C₁-C₁₂haloalkyl, C₂-C₈alkenyl, C₂-C₈alkynyl, C₃-C₈cycloalkyl, -C₁-C₆S(O)₂-R₉, aryl or phenyl-C₁-C₆alkyl; wherein the phenyl- and aryl-radicals may carry from one to three substituents selected independently of one another from nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl and C₁-C₆haloalkoxy;

R₇ is H, OH, C₁-C₁₂alkyl, C₁-C₁₂alkoxy, C₁-C₆alkoxy-C₁-C₆alkoxy, C₂-C₈alkenyloxy, phenyl, phenoxy, benzyloxy, NH₂, NH(C₁-C₁₂alkyl), N(C₁-C₁₂alkyl)₂, -NH-phenyl or -N(C₁-C₁₂alkyl)-phenyl;

R₈ is H, C₁-C₁₂alkyl, phenyl or benzyl; and

R₉ H, C₁-C₁₂alkyl, aryl, aryl-C₁-C₁₂alkyl, or aryl or aryl-C₁-C₁₂alkyl wherein the aryl radical may carry from one to three substituents selected independently of one another from nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl and C₁-C₆haloalkoxy;

and, where appropriate, E/Z isomers, mixtures of E/Z isomers and/or tautomers thereof, in each case in free form or in salt form.

2. A compound according to claim 1 of the formula (I) in free form.
3. A compound according to either of claims 1 or 2 of the formula (I) in which R₁ is isopropyl or sec-butyl.
4. A compound according to either of claims 1 or 3 of the formula (I) in which R₂ is C₁-C₈alkyl.
5. A compound according to either of claims 1 or 4 of the formula (I) in which R₃ is unsubstituted or substituted C₁-C₁₂alkyl.
6. A pesticide composition which comprises at least one compound according to anyone of claims 1 to 5 of the formula (I), in free form or in agrochemically useable salt form, as active ingredient and at least one auxiliary.
7. A process for preparing a composition as described in claim 6, which comprises intimately mixing the active substance with the auxiliary or auxiliaries.
8. A method of controlling pests, which comprises applying a pesticidal composition as described in claim 6 to the pests or their habitat.

9. The use of the compound according to any one of claims 1 to 5 of the formula (I), in free form or, where appropriate, in agrochemically useable salt form, for preparing a composition as described in claim 6.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 02/09315

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 C07H19/01 A01N43/90

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07H A01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 089 202 A (MERCK & CO INC) 21 September 1983 (1983-09-21) the whole document, in particular: page 6, lines 32,33; abstract ---	1-9
X	EP 0 301 806 A (MERCK & CO INC) 1 February 1989 (1989-02-01) page 3, lines 13-16; claim 1 -----	1-3, 5-9

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

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P document published prior to the international filing date but later than the priority date claimed

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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

Fitz, W

INTERNATIONAL SEARCH REPORT

Information on patent family members

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